

## SPECIFICATION

### External Preparation for Skin Diseases Containing Nitroimidazole

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#### Field Of The Invention

The present invention relates to an external preparation which is used for a therapeutics, prophylactic treatment or amelioration of skin diseases which comprises a  
10 nitroimidazole derivative as an active ingredient, a use of the nitroimidazole derivative for producing the external preparation for a therapeutics, prophylactic treatment or amelioration of skin diseases, and the therapeutics and prophylactic treatment of skin diseases using an external  
15 preparation for a therapeutic or prophylactic treatment, or amelioration of skin diseases, which comprises the nitroimidazole derivative as an active ingredient.

#### Background Of The Invention

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##### (Atopic Dermatitis)

An atopic dermatitis is known to be a type I allergic reaction responsive to IgE. Various external preparations for therapeutic treatment of atopic skin disease have been developed so far, which comprises a compound with an activity  
25 for inhibiting PCA reactions which is responsive to IgE, as an active ingredient. However, in a practical application, there has not been known any that is adequately effective, while steroids such as adrenocortical hormones still continue to constitute the mainstream of external preparations for a  
30 therapeutic treatment of atopic skin diseases.

Although the steroids currently used for atopic dermatitis, which are applied for skin diseases and other skin diseases, have excellent therapeutic effects, when used over a long period of time, systemic side effects are  
35 produced including the functional suppression of hypothalamus, pituitary and adrenal cortex. In addition, despite being external preparations, they frequently exhibit local side

effects in the form of skin symptoms such as exacerbation of skin infections and acne characteristic of adrenocortical hormones. Scars, liver spots and freckles in the course of administration as well as problems related to rebound following discontinuation of administration have also been pointed out.

Due to the above problems, immunosuppressants, antihistamines and antiallergics, etc. have been developed as therapeutic agents for atopic dermatitis. However, immunosuppressants involve problems such as exacerbation of bacterial skin infections, and antihistamines involve problems having adverse side effects such as drug rash.

With respect to atopic dermatitis whose cause has not yet be identified, patients having symptoms and their family members thereof suffer from itching and pain on a daily basis and also are perplexed by sleeplessness and various other symptoms. The Patients are not only relying on treatment at hospitals or clinics, but also on private treatment, and etc. Since definitive treatment methods have not yet be established at health care institutes, such as universities, hospitals and etc., there is an urgent need for the development of more effective external preparation for a therapeutic or prophylactic treatment of atopic dermatitis that is free of side effects to take the place of steroid-type anti-inflammatory external preparations.

(Psoriasis)

Psoriasis is one of the most difficult skin diseases to cure and mechanism thereof is unknown. The symptoms recur repeatedly and no fundamental curative method has established so far.

Examples of a therapeutic treatment of psoriasis include application of ointments such as salicylic acid ointment, urea ointment, ointments used for the purpose of moisture retention and vitamin A ointment, heat treatment and soft X-ray, and ointments containing tranilast, cyclosporin or methotrexate, depending on locations and symptoms of the psoriasis. However, these treatments have hardly any

therapeutic efficacy, and external steroid preparations are used mainly for treatment since they are relatively more effective. Today, although the therapeutic efficacy on psoriasis of external steroid preparations is not as great as that for other skin diseases, since there is no other therapeutic treatment available, treatment has been conducted through long-time use of external steroid preparations. However, as widely known, the adverse side effects associated with external steroid preparations are recognized as a problem.

In the treatment of psoriasis, there are problems related to the therapeutic efficacy and adverse side effects of external steroid preparations. Thus, rather than the external steroid preparations, attention has recently been focused on external vitamin D3 preparations. For example, an external active vitamin D3 preparation currently available on the market includes tacalcitol, and this preparation has no adverse side effects as with external steroid preparations, and its therapeutic effects are known to be relatively better than external steroid preparations.

However, even when treated with the above external steroid preparations or external vitamin D3 preparations, the treatment period is normally from several weeks to several months, and there are patients who undergo treatment for a long period of time extending several years to several tens of years. In addition, almost of the patients suffer from relapses which again leads to another period of prolonged treatment.

Thus, if an external preparation were available that could more effectively treat or prevent psoriasis, these types of problems would be able to be solved, therefore, such an external preparation has been desired.

(Hircus, Body Odor and Osmidrosis)

Hircus is the same type of body odor as foot odor, and is thought to occur as a result of components of apocrine perspiration, which is secreted from apocrine glands located in the hair follicles of the skin, being degraded by various

normal flora resulting in the production of a foul odor. As medical treatment of hircus, external preparations such as aluminum chloride solution or formalin alcohol solution are prescribed and used. The treatment is to inhibit odor to a certain degree through antiperspirant action, but they are unable to completely eliminate odor, recurring the odor when perspiration occurs again. In addition, these drugs are also known to cause a high incidence of side effects such as skin rash, itching and rubor following their use.

However, there are currently no medical pharmaceuticals available for the treatment of hircus, foot odor or other forms of body odor in the field of dermatology. Known antiperspirants such as the above aluminum chloride solution and formalin alcohol solution are unable to obtain satisfactory therapeutic results.

Consequently, a surgical operation is employed in which the skin of the armpits is resected to remove the apocrine glands, but this is an extremely bothersome procedure, results in a surgical scar over a wide area following surgery, results in atrophy of skin and muscle, and leaves the skin in a keloid state. Furthermore, there is a considerable economic burden, and relapse is relatively common. In addition, there is frequent occurrence of sequela resulting in neural impairment due to surgical error. The bother, surgical scars and adverse side effects of this surgical treatment along with the mental suffering caused by the sequela are immeasurable for the person undergoing the procedure.

Thus, there has been required for a therapeutic or prophylactic agent for external use for hircus, body odor and osmidrosis that is economical and does not suffer the patient.

(Pigmentation, Blotches and Scars)

For scars due to the sequelae of drug rash, burns, herpes, keloids and smallpox, pigmentation and blotches etc. caused by ultraviolet ray irradiation or cosmetics, there still are no effective external medications despite their having a serious effect on daily life, thereby creating the

need for the development of such a medication.

(Contact Dermatitis, etc.)

There is also a need for an external preparation that is free from side effects and effectively used for a therapeutic or prophylactic treatment of contact dermatitis, plant dermatitis or insect bites, a therapeutic or prophylactic treatment of dermal pruritis or drug rash, a therapeutic or prophylactic treatment of chilblain, a therapeutic or prophylactic treatment of erythroderma, a therapeutic or prophylactic treatment of tinea, a therapeutic or prophylactic treatment of suppurative skin diseases, a therapeutic or prophylactic treatment of pressure sores, a therapeutic or prophylactic treatment of wounds, as well as a therapeutic or prophylactic treatment of palmoplantar pustulosis, lichen planus, lichen nitidus, pityriasis rubra pilaris, pityriasis rosea, erythema (including polymorphic exudative erythema, erythema nodosum and Darier's erythema annulare centrifugum), chronic discoid lupus erythematosus, drug rash and toxic rash, alopecia areata, burns (including scars and keloids), pemphigus, Duhring dermatitis herpetiformis (including pemphigoid), seborrheic dermatitis, dermal stomatitis, Candidiasis (including interdigital erosion, intertrigo, dermal Candidiasis, infantile parasitic erythema, perionychia and vaginal Candidiasis) and tinea versicolor.

It should be noted that the following are known with respect to metronidazole and tinidazole among the nitroimidazole derivatives of the present invention.

A compound metronidazole (2-(2-methyl-5-nitroimidazol-1-yl) ethanol) is a nitroimidazole derivative that was synthesized by Jacob of Rhone-Poulenc Rorer S.A. (France) in 1957. Metronidazole was found to have potent anti-Trichomonas activity by Cosar & Julou. Durel first reported in 1959 that Trichomonas protozoa disappeared following the use of this drug against human trichomoniasis. In addition, it is known that this drug has a strong antimicrobial activity against Entamoeba

histolytica. Moreover, it has also been reported to have bactericidal activity against other anaerobes by both oral administration and topical administration. Its mechanism of action is thought to be the result of the nitro group of metronidazole being reduced by the microorganism, and this reduction in turn leads to a dysfunction such as cleavage in the double stranded DNA of the microorganism thereby inhibiting mitosis and proliferations.

Tinidazole was synthesized in 1966 by the Pfizer Inc. in the United States as a compound having even more potent effect than metronidazole which is an orally used chemotherapeutic agent, and having low toxicity. This compound primarily has anti-Trichomonas activity. Thus, not only does it have superior effects against infections caused by Trichomonas vaginalis as well as against Trichomonas vaginalis infecting the vulva, cervical tube, urinary tract and rectum, but also exhibits antimicrobial activity against anaerobic microorganisms as well and is used clinically for such purposes. Its mechanism of action is thought to involve reduction of the nitro group of tinidazole by the microorganism resulting in this reduction product causing functional impairment such as cleavage of double strand DNA, thereby inhibiting mitosis and proliferations of the microorganism.

In addition, with respect to metronidazole, the following information is known regarding the effect of its administration on immunity. Namely, it is clearly indicated in Int. Arch. Allergy Appl. Immun., 54, 422 (1977) that, in mice orally administered metronidazole, although the formation of granuloma by intravenous injection of the eggs of Schistosoma mansoni was inhibited, non-specific granuloma formation was not inhibited. According to Int. J. Radiation Oncology Biol Phys., 2, 701 (1983), intraperitoneal administration of metronidazole is known to inhibit swelling of the ears induced by dinitrofluorobenzene in mice sensitized with dinitrofluorobenzene. In addition, according to the Indian J. Exp. Biol., 25, 177 (1987), it is clearly

indicated that intraperitoneal administration of metronidazole significantly inhibits increases in anti-TBA antibody titer to TBA vaccine in rabbits, and according to Indian J. Exp. Biol., 29, 867 (1991), it is clearly indicated that intraperitoneal administration of metronidazole inhibits a delayed immune reaction to intravenous injection of ovine erythrocytes while also demonstrating inhibitory action on leukocyte migration. Moreover, known effects of metronidazole on inflammation include metronidazole external preparations being effective against inflammatory skin diseases such as rosacea (International Unexamined Patent Publication No. WO88/06888, International Unexamined Patent Publication No. WO89/06537, International Unexamined Patent Publication No. WO94/08350, International Unexamined Patent Publication No. WO96/01117 and International Unexamined Patent Publication No. WO98/27960). In addition, according to Mykosen, 27, 475 (1984), the reason why metronidazole exhibits therapeutic effects at a concentration at which it does not exhibit antimicrobial activity against P. ovale and so forth is because of its anti-inflammatory activity, and according to Br. J. Dermatol., 114, 231 (1986), metronidazole has inhibitory activity on the production of active oxygen species, and the reason why metronidazole is effective against rosacea is partially due to its anti-inflammatory activity. International Surgery, 60, 75 (1975) indicates that oral administration of metronidazole is effective against pododermal ulcers.

On the other hand, with respect to tinidazole, it is described in Indian J. Exp. Biol., 29, 867 (1991) with respect to immunity that intraperitoneal administration of tinidazole tends to inhibit delayed immune reaction to intravenous injection of ovine erythrocytes, and that it clearly exhibits inhibitory action of leukocyte migration. Moreover, with respect to inflammation, tinidazole external preparations are known to be used for the therapeutic treatment of skin inflammation (International Unexamined Patent Publication No. WO93/20817, International Unexamined

Patent Publication No. WO98/27960).

With respect to the use of metronidazole for the therapeutic treatment of psoriasis, it is disclosed in US Patent Publication No. US 4,491,588 that oral preparations of metronidazole are effective for the treatment of psoriasis, and in International Unexamined Patent Publication No. WO96/01117, psoriasis is indicated as being one of the inflammatory diseases that can be cured with external preparations of metronidazole.

However, in those references mentioned above pertaining to immunity, with the exception of the Int. J. Radiation Oncology Biol. Phys., 2, 701 (1983), all of the immune reactions observed are immune reactions other than on the skin surface. In addition, the observed immunosuppressive effects are remarkably lower in comparison with those of immunosuppressants used clinically, and it is therefore considered that external preparations of metronidazole or tinidazole cannot be expected to be effective as therapeutic agents for atopic dermatitis. Further, there is no correlation between the effectiveness in treatment of atopic dermatitis and the effectiveness in the model of contact dermatitis used in the Int. J. Radiation Oncology Biol. Phys., 2, 701 (1983) in which the only immune reaction on the skin surface is observed. Moreover, it has not been known that typical therapeutic agents for inflammatory disease are used for therapeutic treatment of atopic dermatitis. In addition, the use of a nitroimidazole derivative for treatment of atopic dermatitis is also previously unknown.

Further, US Patent No. 4,491,588 discloses the treatment of psoriasis by oral administration of metronidazole, and although ketoconazole, which is similarly disclosed as being effective in the treatment of psoriasis, has been granted a right as an oral preparation (US Patent No. 4,491,588) and as an external preparation (US Patent No. 4,569,935), only an oral preparation has been granted a right with respect to metronidazole. The present invention is directed to findings that an external preparation of metronidazole is superior to

the oral preparation in terms of effect and toxicity.  
Moreover, since the therapeutic use for psoriasis indicated  
in International Unexamined Patent Publication No. WO96/01117  
is one example of a typical inflammatory disease, and the  
5 disclosed contents merely indicate that an external  
preparation of metronidazole is able to inhibit the formation  
of edema caused by local stimulation by arachidonic acid, as  
was described by the applicant himself to the effect that,  
"conventional non-steroid anti-inflammatory drugs, such as  
10 cyclooxygenase or lipoxygenase reaction inhibitors (including  
indometacin, naproxen and phenylbutazone) and preparations  
able to inhibit conduit plasma backflow (such as  
vasoconstrictors) are excellent reaction inhibitors in this  
model", this is an experimental system in which conventional  
15 non-steroid anti-inflammatory drugs (NSAIDs) also exhibit  
excellent effect. In this publication, it is deduced that  
metronidazole can be used for the treatment of "eczema,  
psoriasis, rosacea, lupus vulgaris, ulcers and seborrheic  
dermatitis", etc. only by virtue of confirming its action.  
20 However, this patent application cannot be included in a  
prior art reference of the present application since the  
etiology of psoriasis is unknown, nearly all NSAIDs do not  
exhibit therapeutic effects against psoriasis and the  
therapeutic effect against psoriasis has actually not been  
25 confirmed.

#### Disclosure of the Invention

The present inventors have made extensive and intensive  
studies on a therapeutic or prophylactic agent for atopic  
30 dermatitis, and have found that an external preparation  
containing a nitroimidazole derivative as an active  
ingredient is extremely effective as a therapeutic or  
prophylactic agent for atopic dermatitis, and have found that  
the invention is highly safe and is free from adverse side  
35 effects, and thus, the present invention has been completed.  
In addition, the present inventors also have found that it is  
also particularly effective for a therapeutic and

prophylactic treatment of facial atopy and pediatric atopy, for which the treatment has been difficult.

Moreover, the present inventors have found that an external preparation containing a nitroimidazole derivative  
5 is effective for ameliorating blotches, pigmentation or scarring of the skin, effective for a therapeutic or prophylactic treatment of psoriasis, and effective for a therapeutic or prophylactic treatment of hircus, body odor or osmidrosis. In addition, the present inventors have also  
10 found that an external preparation containing a nitroimidazole derivative is effective for a therapeutic or prophylactic treatment of contact dermatitis, plant dermatitis or insect bites, a therapeutic or prophylactic treatment of dermal pruritis or drug rash, a therapeutic or  
15 prophylactic treatment of chilblain, a therapeutic or prophylactic treatment of erythroderma, a therapeutic or prophylactic treatment of tinea, a therapeutic or prophylactic treatment of suppurative skin diseases, a therapeutic or prophylactic treatment of pressure sores, a  
20 therapeutic or prophylactic treatment of wounds, and a therapeutic or prophylactic treatment of palmoplantar pustulosis, lichen planus, lichen nitidus, pityriasis rubra pilaris, pityriasis rosea, erythema (including polymorphic exudative erythema, erythema nodosum and Darier's erythema  
25 annulare centrifugum), chronic discoid lupus erythematosus, drug rash and toxic rash, alopecia areata, burns (including scars and keloids), pemphigus, Duhring dermatitis herpetiformis (including pemphigoid), seborrheic dermatitis, dermal stomatitis, Candidiasis (including interdigital  
30 erosion, intertrigo, dermal Candidiasis, infantile parasitic erythema, perionychia and vaginal Candidiasis) or tinea versicolor.

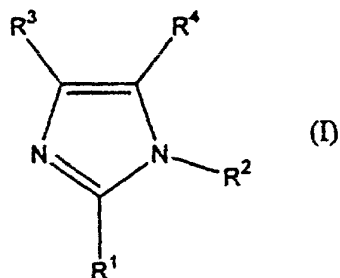
The present inventors have found that antiamebic effects appear rapidly when crotamiton is contained in an  
35 external preparation containing a nitroimidazole derivative.

Other features of the present invention include a use of the nitroimidazole derivative for producing of an external

preparation for skin disease for a prophylactic or  
therapeutic treatment of atopic dermatitis, amelioration of  
skin blotches, pigmentation or scarring, a therapeutic or  
prophylactic treatment of psoriasis, and a therapeutic or  
5 prophylactic treatment of hircus, body odor or osmidrosis, as  
well as a therapeutic or prophylactic treatment and  
amelioration of these diseases using an external preparation  
for skin diseases containing the nitroimidazole derivative.

In addition, the present inventors also have found that  
10 an external preparation which comprises at least one compound  
of the nitroimidazole derivatives and at least one agent  
selected from the group consisting of an antimycotic agent,  
antibacterial agent, sulfa, immunosuppressant, anti-  
inflammatory agent, antibiotic, antiviral agent, metabolic  
15 antagonist, antihistamine, tissue repair promoter, vitamin,  
antiallergic, local anesthetic, hair agent and steroid being  
administered simultaneously or separately with an interval  
allows a concentration of these drugs other than  
nitroimidazole to be reduced, thereby eliminating adverse  
20 side effects while also being fast-acting. Moreover, it was  
also found that similar effects are demonstrated even at  
concentrations at which these agents other than  
nitroimidazole do not exhibit pharmacological effects.

The external preparation for a therapeutic or  
25 prophylactic treatment or amelioration of skin disease  
according to the present invention comprises a nitroimidazole  
derivative represented by the following formula (I), a  
pharmaceutically acceptable salt thereof, an ester thereof or  
other derivatives thereof as an active ingredient:



30 wherein R<sup>1</sup>, R<sup>3</sup> and R<sup>4</sup> may be the same or different from each

other and each independently represent a hydrogen atom, a nitro group, a lower alkyl group, a lower alkyl group substituted by 1 or more substituents which may be the same or different selected from a Substituent group  $\alpha$  and

5 Substituent group  $\beta$ , a lower alkenyl group or a lower alkenyl group substituted by 1 or more substituents which may be the same or different selected from the Substituent group  $\alpha$  and the Substituent group  $\beta$ ; and  $R^2$  represents a hydrogen atom, a lower alkyl group, a lower alkyl group substituted by 1 or

10 more substituents which may be the same or different selected from the Substituent group  $\alpha$  and the Substituent group  $\beta$ , a lower alkenyl group or a lower alkenyl group substituted by 1 or more substituents which may be the same or different selected from the Substituent group  $\alpha$  and the Substituent

15 group  $\beta$ , provided that any one of  $R^1$ ,  $R^3$  and  $R^4$  is a nitro group,

Substituent group  $\alpha$

comprises a lower alkyloxy group, a lower alkyloxy group substituted by 1 or more substituents which may be the same

20 or different selected from the Substituent group  $\beta$ , a lower alkylcarbonyloxy group, a lower alkylcarbonyloxy group substituted by 1 or 2 or more substituents which may be the same or different selected from the Substituent group  $\beta$ , a lower alkylsulfonyl group, a lower alkylsulfonyl group

25 substituted by 1 or more substituents which may be the same or different selected from the Substituent group  $\beta$ , a cycloalkyl group, a cycloalkyl group substituted by 1 or more substituents which may be the same or different selected from the Substituent group  $\beta$ , a heteroaryl group, a heteroaryl

30 group substituted by 1 or more substituents which may be the same or different selected from the Substituent group  $\beta$ , an aryl group and an aryl group substituted by 1 or 2 or more substituents which may be the same or different selected from the Substituent group  $\beta$ .

35 Substituent group  $\beta$

comprises a hydroxy group, a mercapto group, a halogen atom,

an amino group, a lower alkylamino group, a lower alkyloxy group, a lower alkenyl group, a cyano group, a carboxy group, a carbamoyloxy group, a carboxyamide group, a thiocarboxyamide group a morpholino group and etc.

5 Of the above external preparations, preferred are

(1) an external preparation wherein  $R^4$  is a nitro group,  
(2) an external preparation of (1) above wherein  $R^1$  and  $R^2$   
are the same or different and represent a lower alkyl group,  
a lower alkyl group substituted by 1 or 2 or more

10 substituents selected from the Substituent group  $\alpha$  and the  
Substituent group  $\beta$ , a lower alkenyl group, or a lower  
alkenyl group substituted by 1 or 2 or more substituents  
which may be the same or different selected from the  
Substituent group  $\alpha$  and the Substituent group  $\beta$ , and  $R^3$  is a  
15 hydrogen atom,

(3) an external preparation of (2) wherein <Substituent group  
 $\alpha$ > is a lower alkyloxy group and the Substituent group  $\beta$  is a  
hydroxy group, an amino group, a halogen atom, a cycloalkyl  
group, a heteroaryl group or an aryl group,

20 (4) an external preparation of (3) wherein the Substituent  
group  $\beta$  is a hydroxy group, an amino group, a halogen atom or  
a heteroaryl group,

(5) an external preparation of (3) wherein  $R^1$  is a lower  
alkyl group,

25 (6) an external preparation of (3) wherein  $R^2$  is a lower  
alkyl group substituted by a hydroxy group,

(7) an external preparation of (2) wherein the Substituent  
group  $\alpha$  is a lower alkylsulfonyl group or a lower  
alkylsulfonyl group substituted by substituents which may be

30 the same or different selected from the Substituent group  $\beta$   
and the Substituent group  $\beta$  is a hydroxy group, a halogen  
atom, an amino group, a lower alkylamino group, a lower  
alkyloxy group, a lower alkenyl group, a cyano group, a  
carboxy group, a cycloalkyl group or an aryl group and etc,

35 (8) an external preparation of (7) wherein  $R^1$  is a lower  
alkyl group or a lower alkyl group substituted by substituent

which may be the same or different selected from Substituent group  $\beta$ , and

(9) an external preparation of (7) wherein  $R^2$  is a lower alkylsulfonyl group or a lower alkyl group substituted by the lower alkylsulfonyl group substituted by substituent which may be the same or different selected from the Substituent group  $\beta$ .

The above (1) and (2), (3) to (5) or (7) and (8) represent more preferable compound as the number becomes larger. In the general formula (I), external preparations obtained by optionally selecting  $R^1$  to  $R^4$  from (1) to (9) and optionally combining those are also preferable and the more preferable external preparations are (5)-(6) and (8)-(9). Still further preferable external preparations are selected from the following groups:

Compound group

2-(2-methyl-5-nitroimidazol-1-yl)ethanol (general name: metronidazole) and, 1-(2-ethylsulfonyl-ethyl)-2-methyl-5-nitroimidazole (general name: tinidazole).

In the above, examples of the "lower alkyl group" of  $R^1$  to  $R^4$  and the "lower alkyl group" of the "lower alkyl group substituted by 1 or more substituents which may be the same or different selected from the Substituent group  $\alpha$  and the Substituent group  $\beta$ " may include a straight or branched alkyl group having 1 to 6 carbon atoms such as methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, s-butyl, tert-butyl, n-pentyl, isopentyl, 2-methylbutyl, neopentyl, 1-ethylpropyl, n-hexyl, isohexyl, 4-methylpentyl, 3-methylpentyl, 2-methylpentyl, 1-methylpentyl, 3,3-dimethylbutyl, 2,2-dimethylbutyl, 1,1-dimethylbutyl, 1,2-dimethylbutyl, 1,3-dimethylbutyl, 2,3-dimethylbutyl and 2-ethylbutyl, preferably the straight or branched alkyl group having 1 to 3 carbon atoms, more preferably the methyl group in  $R^1$  and the ethyl group in  $R^2$ .

In the above formula, examples of the "lower alkenyl group" in  $R^1$  to  $R^4$  and Substituent group  $\alpha$  and Substituent group  $\beta$  and the "lower alkenyl group" of the "lower alkenyl

group substituted by 1 or 2 or more substituents which may be the same or different selected from the Substituent group  $\alpha$  and the Substituent group  $\beta$ " may include a straight or branched alkenyl group having 2 to 6 carbon atoms such as ethenyl, 1-propenyl, 2-propenyl, 1-methyl-2-propenyl, 1-methyl-1-propenyl, 2-methyl-1-propenyl, 2-methyl-2-propenyl, 2-ethyl-2-propenyl, 1-butenyl, 2-butenyl, 1-methyl-2-butenyl, 1-methyl-1-butenyl, 3-methyl-2-butenyl, 1-ethyl-2-butenyl, 3-butenyl, 1-methyl-3-butenyl, 2-methyl-3-butenyl, 1-ethyl-3-butenyl, 1-pentenyl, 2-pentenyl, 1-methyl-2-pentenyl, 2-methyl-2-pentenyl, 3-pentenyl, 1-methyl-3-pentenyl, 2-methyl-3-pentenyl, 4-pentenyl, 1-methyl-4-pentenyl, 2-methyl-4-pentenyl, 1-hexenyl, 2-hexenyl, 3-hexenyl, 4-hexenyl 5-hexenyl and etc., preferably the straight or branched alkenyl group having 3 to 5 carbon atoms.

In the above formula (I), examples of the "halogen atom" in Substituent group  $\beta$  may include a fluorine atom, a chlorine atom, a bromine atom and an iodine atom, preferably a fluorine atom and a chlorine atom.

In the above formula (I), the "lower alkyloxy group" in the Substituent group  $\alpha$  and the Substituent group  $\beta$  and the "lower alkyloxy group" of the "lower alkyloxy group" substituted by 1 or more substituents which may be the same or different selected from the Substituent group  $\beta$ " represent a group in which the above-mentioned "lower alkyl group" is bonded to an oxygen atom and examples of such group may include a straight or branched alkyloxy group having 1 to 6 carbon atoms such as methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, isobutoxy, s-butoxy, tert-butoxy, n-pentoxy, isopentoxy, 2-methylbutoxy, neopentoxy, n-hexyloxy, 4-methylpentoxy, 3-methylpentoxy, 2-methylpentoxy, 3,3-dimethylbutoxy, 2,2-dimethylbutoxy, 1,1-dimethylbutoxy, 1,2-dimethylbutoxy, 1,3-dimethylbutoxy, 2,3-dimethylbutoxy and etc., preferably a straight or branched alkyloxy group having 1 to 3 carbon atoms, more preferably a methoxy group.

In the above formula (I), the "lower alkylcarbonyloxy group" in Substituent group  $\alpha$  and the "lower alkylcarbonyloxy group" in Substituent group  $\beta$  may include a straight or branched alkylcarbonyloxy group having 1 to 6 carbon atoms, preferably a straight or branched alkylcarbonyloxy group having 1 to 3 carbon atoms, more preferably a formyl group.

group" of the "lower alkylcarbonyloxy group substituted by 1 or more substituents which may be the same or different selected from the Substituent group  $\beta$ " represent a group in which the above "lower alkyl group" is bonded to a

carbonyloxy group and examples of such group may include a straight or branched alkylcarbonyloxy group having 2 to 7 carbon atoms such as acetyloxy, propionyloxy, butyryloxy, isobutyryloxy, pentanoyloxy, pivaloyloxy, valeryloxy, isovaleryloxy and hexanoyloxy, preferably a straight or branched alkylcarbonyloxy group having 2 to 4 carbon atoms, more preferably a formyloxy group or an acetyloxy group.

In the above formula (I), the "lower alkylsulfonyl group" in Substituent group  $\alpha$  and the "lower alkylsulfonyl group" of the "lower alkylsulfonyl group substituted by 1 or 2 or more substituents which may be the same or different selected from <Substituent group  $\beta$ >" represent a group in which the above "lower alkyl group" is bonded to a sulfonyl group and examples of such group may include a straight or branched alkylsulfonyl group having 1 to 6 carbon atoms such as methanesulfonyl, ethanesulfonyl, n-propanesulfonyl, isopropanesulfonyl, n-butanesulfonyl, isobutanesulfonyl, s-butanesulfonyl, tert-butanesulfonyl, n-pentanesulfonyl, isopentanesulfonyl, 2-methylbutanesulfonyl, neopentanesulfonyl, n-hexanesulfonyl, 4-methylpentanesulfonyl, 3-methylpentanesulfonyl, 2-methylpentanesulfonyl, 3,3-dimethylbutanesulfonyl, 2,2-dimethylbutanesulfonyl, 1,1-dimethylbutanesulfonyl, 1,2-dimethylbutanesulfonyl, 1,3-dimethylbutanesulfonyl, 2,3-dimethylbutanesulfonyl and etc., preferably a straight or branched alkylsulfonyl group having 1 to 3 carbon atoms, more preferably an ethanesulfonyl group.

In the above formula (I), the "lower alkylamino group" in Substituent group  $\beta$  represents a group in which the above "lower alkyl group" is substituted by an amino group and examples of such group may include a straight or branched alkylamino group having 1 to 6 carbon atoms such as methylamino, ethylamino, n-propylamino, isopropylamino, n-butylamino, isobutylamino, s-butylamino, tert-butylamino, n-

pentylamino, isopentylamino, 2-methylbutylamino, neopentylamino, 1-ethylpropylamino, n-hexylamino, isohexylamino, 4-methylpentylamino, 3-methylpentylamino, 2-methylpentylamino, 1-methylpentylamino, 3,3-dimethylbutylamino, 2,2-dimethylbutylamino, 1,1-dimethylbutylamino, 1,2-dimethylbutylamino, 1,3-dimethylbutylamino, 2,3-dimethylbutylamino, 2-ethylbutylamino and etc., preferably a straight or branched alkylamino group having 1 to 6 carbon atoms, more preferably a methylamino group or an ethylamino group.

In the above formula (I), the "cycloalkyl group" in Substituent group  $\alpha$  and the "cycloalkyl group" of the "cycloalkyl group substituted by 1 or more substituents which may be the same or different selected from the Substituent group  $\beta$ " may include a 3 to 10-membered saturated cyclic hydrocarbon group which may be condensed such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, norbornyl and adamantyl, preferably a 5 to 7-membered saturated cyclic hydrocarbon group.

In the above formula (I), examples of the "heteroaryl group" in Substituent group  $\alpha$  and the "heteroaryl group" of the "heteroaryl group substituted by 1 or more substituents which may be the same or different selected from the Substituent group  $\beta$ " may include a 5 to 7-membered aromatic heterocyclic group such as furyl, thienyl, pyrrolyl, azepinyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, 1,2,3-oxadiazolyl, triazolyl, tetrazolyl, thiadiazolyl, pyranyl, pyridyl, pyridazinyl, pyrimidinyl, pyrazinyl and etc., preferably a pyridyl group.

In the above formula (I), examples of the "aryl group" in Substituent group  $\alpha$  and the "aryl group" of the "aryl group substituted by 1 or more substituents which may be the same or different selected from the Substituent group  $\beta$ " may include an aromatic hydrocarbon group having 5 to 14 carbon atoms such as phenyl, indenyl, naphthyl, phenanthrenyl, anthrathenyl and etc., preferably an aromatic hydrocarbon group having 6 to 10 carbon atoms, more preferably a phenyl

group.

Since the compound (I) of the present invention can be converted to a salt, the "pharmacologically acceptable salt thereof" represents such salt and may preferably include

5 metal salts such as alkali metal salts, e.g., a sodium salt, a potassium salt and a lithium salt and etc., metal salt such as alkaline earth metal salts, e.g., a calcium salt and a magnesium salt, an aluminum salt, an iron salt, a zinc salt, a copper salt, a nickel salt and a cobalt salt; amine salts  
10 such as inorganic salts, e.g., an ammonium salt and organic salts, e.g., a t-octylamine salt, a dibenzylamine salt, a morpholine salt, a glucosamine salt, a phenylglycinealkyl ester salt, an ethylenediamine salt, an N-methylglucamine salt, a guanidine salt, a diethylamine salt, a triethylamine  
15 salt, a dicyclohexylamine salt, an N,N'-dibenzylethylenediamine salt, a chloroprocain salt, a procain salt, a diethanolamine salt, an N-benzyl-phenethylamine salt, a piperazine salt, a tetramethylammonium salt, a tris(hydroxymethyl)aminomethane salt and etc.; inorganic acid  
20 salts such as halogenated hydroacid salts, e.g., a hydrofluoric acid salt, a hydrochloric acid salt, a hydrobromic acid salt and a hydroiodic acid salt, a nitric acid salt, a perchloric acid salt, a sulfuric acid salt, a phosphoric acid salt and etc.; organic acid salts such as  
25 lower alkanesulfonic acid salt, e.g., a methanesulfonic acid salt, a trifluoromethanesulfonic acid salt and an ethanesulfonic acid salt, arylsulfonic acid salts, e.g., a benzenesulfonic acid salt and a p-toluenesulfonic acid salt, an acetic acid salt, a malic acid salt, a fumaric acid salt,  
30 a succinic acid salt, a citric acid salt, a tartaric acid salt, an oxalic acid salt and maleic acid salt; and amino acid salts such as a glycine salt, a lysine salt, an arginine salt, an ornithine salt, a glutamic acid salt and an aspartic acid salt. In the case where the salt becomes the metal  
35 salts or the amine salts, it is limited to the case where the compound (I) has an acidic group.

Moreover, in case the compound (I) of the present

invention absorbs moisture content by allowing it to stand in the atmosphere so that adsorbed water is deposited thereon to form a hydrate, such salt is also included in the present invention.

5 Further, in case the compound (I) of the present invention absorbs a certain kind of solvent to form a solvate, such salt is also included in the present invention.

Since the compound (I) of the present invention can be converted to an ester, the "ester thereof" means such ester and represents "ester of a hydroxy group" and "ester of a carboxy group", wherein the respective ester residues are "a general protective group" or "a protective group cleavable by a biological method such as hydrolysis in a living body".

10 The "general protective group" represents a protective group cleavable by a chemical method such as hydrogenation decomposition, hydrolysis, electrolysis and optical decomposition and the "general protective group" relating to the "ester of the hydroxy group" may include "an aliphatic acyl group" such as an alkylcarbonyl group, e.g., formyl, acetyl, propionyl, butyryl, isobutyryl, pentanoyl, pivaloyl, valeryl, isovaleryl, octanoyl, nonanoyl, decanoyl, 3-methylnonanoyl, 8-methylnonanoyl, 3-ethyloctanoyl, 3,7-dimethyloctanoyl, undecanoyl, dodecanoyl, tridecanoyl, tetradecanoyl, pentadecanoyl, hexadecanoyl, 1-methylpentadecanoyl, 14-methylpentadecanoyl, 13,13-dimethyltetradecanoyl, heptadecanoyl, 15-methylhexadecanoyl, octadecanoyl, 1-methylheptadecanoyl, nonadecanoyl, eicosanoyl, heneicosanoyl and etc., a carboxylated alkylcarbonyl group, e.g., succinoyl, glutanoyl and adipoyl, a halogeno-lower alkylcarbonyl group, e.g., chloroacetyl, dichloroacetyl, trichloroacetyl and trifluoroacetyl, a saturated cyclic hydrocarbon-carbonyl group, e.g., cyclopropylcarbonyl, cyclobutylcarbonyl, cyclopentylcarbonyl, cyclohexylcarbonyl, cycloheptylcarbonyl and cyclooctylcarbonyl, a lower alkoxy lower alkylcarbonyl group, e.g., methoxyacetyl, and an unsaturated alkylcarbonyl group, e.g., (E)-2-methyl-2-butenoyl;

"an aromatic acyl group" such as an arylcarbonyl group, e.g., benzoyl,  $\alpha$ -naphthoyl,  $\beta$ -naphthoyl, pyridoyl, thienoyl and furoyl, a halogeno arylcarbonyl group, e.g., 2-bromobenzoyl and 4-chlorobenzoyl, a lower alkylated arylcarbonyl group, e.g., 2,4,6-trimethylbenzoyl and 4-toluoyl, a lower alkoxyated arylcarbonyl group, e.g., 4-anisoyl, a carboxylated arylcarbonyl group, e.g., 2-carboxybenzoyl, 3-carboxybenzoyl and 4-carboxybenzoyl, a nitrated arylcarbonyl group, e.g., 4-nitrobenzoyl and 2-nitrobenzoyl, a lower alkoxy carbonylated arylcarbonyl group, e.g., 2-(methoxycarbonyl)benzoyl an arylated arylcarbonyl group and etc, e.g., 4-phenylbenzoyl; "an aralkylcarbonyl group" such as a lower alkylcarbonyl group substituted by 1 to 3 aryl groups, e.g., phenylacetyl,  $\alpha$ -naphthylpropionyl,  $\beta$ -naphthylbutyryl, diphenylisobutyryl, triphenylacetyl,  $\alpha$ -naphthylidiphenylisobutyryl and 9-anthrylpentanoyl and a lower alkylcarbonyl group substituted by 1 to 3 aryl groups of which aryl ring is substituted by a lower alkyl group, a lower alkoxy group, a nitro group, a halogen atom, a cyano group and etc., e.g., 4-methylphenylacetyl, 2,4,6-trimethylphenylformyl, 3,4,5-trimethylphenylbutyryl, 4-methoxyphenylisobutyryl, 4-methoxyphenyldiphenylpivaloyl, 2-nitrophenylacetyl, 4-nitrophenylpropionyl, 4-chlorophenylbutyryl, 4-bromophenylacetyl and 4-cyanophenylpentanoyl; "a tetrahydropyranyl group or a tetrahydrothiopyranyl group" such as tetrahydropyran-2-yl, 3-bromotetrahydropyran-2-yl, 4-methoxytetrahydropyran-4-yl, tetrahydrothiopyran-2-yl and 4-methoxytetrahydrothiopyran-4-yl; "a tetrahydrofuran-yl group or a tetrahydrothiofuran-yl group" such as tetrahydrofuran-2-yl and tetrahydrothiofuran-2-yl; "a silyl group" such as a tri-lower alkylsilyl group, e.g., trimethylsilyl, triethylsilyl, isopropyldimethylsilyl, t-butyldimethylsilyl, methyldiisopropylsilyl, methyldi-t-butylsilyl and triisopropylsilyl and a tri-lower alkylsilyl group substituted by 1 or 2 aryl groups, e.g., diphenylmethylsilyl, diphenylbutylsilyl, diphenylisopropylsilyl and phenyldiisopropylsilyl; "an alkoxymethyl group" such as a

lower alkoxymethyl group, e.g., methoxymethyl, 1,1-dimethyl-1-methoxymethyl, ethoxymethyl, propoxymethyl, isopropoxymethyl, butoxymethyl and tert-butoxymethyl, a lower alkoxylated lower alkoxymethyl group, e.g., 2-

5 methoxyethoxymethyl and

a halogeno-lower alkoxymethyl group, e.g., 2,2,2-

trichloroethoxymethyl and bis(2-chloroethoxy)methyl;

"a substituted ethyl group" such as a lower alkoxylated ethyl group, e.g., 1-ethoxyethyl and 1-(isopropoxy)ethyl and a

10 halogenated ethyl group, e.g., 2,2,2-trichloroethyl; "an

aralkyl group" such as a lower alkyl group substituted by 1

to 3 aryl groups, e.g., benzyl,  $\alpha$ -naphthylmethyl,  $\beta$ -

naphthylmethyl, diphenylmethyl, triphenylmethyl,  $\alpha$ -

naphthyldiphenylmethyl and 9-anthrylmethyl and a lower alkyl

15 group substituted by 1 to 3 aryl groups of which aryl ring is substituted by a lower alkyl group, a lower alkoxy group, a

nitro group, a halogen atom or a cyano group, e.g., 4-

methylbenzyl, 2,4,6-trimethylbenzyl, 3,4,5-trimethylbenzyl,

4-methoxybenzyl, 4-methoxyphenyldiphenylmethyl, 2-nitrobenzyl,

20 4-nitrobenzyl, 4-chlorobenzyl, 4-bromobenzyl and 4-

cyanobenzyl; "an alkoxycarbonyl group" such as a lower

alkoxycarbonyl group, e.g., methoxycarbonyl, ethoxycarbonyl,

tert-butoxycarbonyl and isobutoxycarbonyl and a lower

alkoxycarbonyl group substituted by a halogen atom or a tri-

25 lower alkylsilyl group, e.g., 2,2,2-trichloroethoxycarbonyl,

2-trimethylsilylethoxycarbonyl and etc.; "an

alkenyloxycarbonyl group" such as vinyloxycarbonyl and

allyloxycarbonyl; and "an aralkyloxycarbonyl group" of which

aryl ring may be substituted by 1 or 2 lower alkoxy or nitro

30 groups such as benzyloxycarbonyl, 4-methoxybenzyloxycarbonyl,

3,4-dimethoxybenzyloxycarbonyl, 2-nitrobenzyloxycarbonyl and

4-nitrobenzyloxycarbonyl and etc.

Whereas, the "general protective group" relating to the

"ester of the carboxy group" may preferably include "a lower

35 alkyl group" such as methyl, ethyl, n-propyl, isopropyl, n-

butyl, isobutyl, s-butyl, tert-butyl, n-pentyl, isopentyl, 2-

methylbutyl, neopentyl, 1-ethylpropyl, n-hexyl, isohexyl, 4-

methylpentyl, 3-methylpentyl, 2-methylpentyl, 1-methylpentyl,  
 3,3-dimethylbutyl, 2,2-dimethylbutyl, 1,1-dimethylbutyl, 1,2-  
 dimethylbutyl, 1,3-dimethylbutyl, 2,3-dimethylbutyl, 2-  
 ethylbutyl and etc.; "an alkenyl group" such as ethenyl, 1-  
 5 propenyl, 2-propenyl, 1-methyl-2-propenyl, 1-methyl-1-  
 propenyl, 2-methyl-1-propenyl, 2-methyl-2-propenyl, 2-ethyl-  
 2-propenyl, 1-butenyl, 2-butenyl, 1-methyl-2-butenyl, 1-  
 methyl-1-butenyl, 3-methyl-2-butenyl, 1-ethyl-2-butenyl, 3-  
 butenyl, 1-methyl-3-butenyl, 2-methyl-3-butenyl, 1-ethyl-3-  
 10 butenyl, 1-pentenyl, 2-pentenyl, 1-methyl-2-pentenyl, 2-  
 methyl-2-pentenyl, 3-pentenyl, 1-methyl-3-pentenyl, 2-methyl-  
 3-pentenyl, 4-pentenyl, 1-methyl-4-pentenyl, 2-methyl-4-  
 pentenyl, 1-hexenyl, 2-hexenyl, 3-hexenyl, 4-hexenyl, 5-  
 hexenyl and etc.; "an alkynyl group" such as ethynyl, 2-  
 15 propynyl, 1-methyl-2-propynyl, 2-methyl-2-propynyl, 2-ethyl-  
 2-propynyl, 2-butyne, 1-methyl-2-butyne, 2-methyl-2-butyne,  
 1-ethyl-2-butyne, 3-butyne, 1-methyl-3-butyne, 2-methyl-3-  
 butynyl, 1-ethyl-3-butyne, 2-pentyne, 1-methyl-2-pentyne,  
 2-methyl-2-pentyne, 3-pentyne, 1-methyl-3-pentyne, 2-  
 20 methyl-3-pentyne, 4-pentyne, 1-methyl-4-pentyne, 2-methyl-  
 4-pentyne, 2-hexynyl, 3-hexynyl, 4-hexynyl, 5-hexynyl and  
 etc.; "a halogeno- lower alkyl group" such as trifluoromethyl,  
 trichloromethyl, difluoromethyl, dichloromethyl,  
 dibromomethyl, fluoromethyl, 2,2,2-trifluoroethyl, 2,2,2-  
 25 trichloroethyl, 2-bromoethyl, 2-chloroethyl, 2-fluoroethyl,  
 2-iodoethyl, 3-chloropropyl, 4-fluorobutyl, 6-iodohexyl and  
 2,2-dibromoethyl; "a hydroxy lower alkyl group" such as 2-  
 hydroxyethyl, 2,3-dihydroxypropyl, 3-hydroxypropyl, 3,4-  
 dihydroxybutyl 4-hydroxybutyl and etc.; "an aliphatic acyl"-  
 30 "lower alkyl group" such as acetylmethyl; "an aralkyl group"  
 such as "a lower alkyl group" substituted by 1 to 3 aryl  
 groups, e.g., benzyl, phenethyl, 3-phenylpropyl,  $\alpha$ -  
 naphthylmethyl,  $\beta$ -naphthylmethyl, diphenylmethyl,  
 triphenylmethyl, 6-phenylhexyl,  $\alpha$ -naphthylmethyl and  
 35 9-anthrylmethyl and etc., and a lower alkyl group substituted  
 by 1 to 3 aryl group of which aryl ring is substituted by a  
 lower alkyl group, a lower alkoxy group, a nitro group, a

halogen atom, a cyano group or an alkoxy carbonyl group, e.g.,  
4-methylbenzyl, 2,4,6-trimethylbenzyl, 3,4,5-trimethylbenzyl,  
4-methoxybenzyl, 4-methoxyphenyldiphenylmethyl, 2-nitrobenzyl,  
4-nitrobenzyl, 4-chlorobenzyl, 4-bromobenzyl, 4-cyanobenzyl,  
5 4-cyanobenzoyldiphenylmethyl, bis(2-nitrophenyl)methyl,  
piperonyl, 4-methoxycarbonylbenzyl and etc.; and "a silyl  
group" such as trimethylsilyl, triethylsilyl,  
isopropyldimethylsilyl, tert-butyldimethylsilyl,  
methyldiisopropylsilyl, methyldi-tert-butylysilyl,  
10 triisopropylsilyl, methyldiphenylsilyl,  
isopropyldiphenylsilyl, butyldiphenylsilyl,  
phenyldiisopropylsilyl, and etc.

"The protective group cleavable by a biological method  
such as hydrolysis in a living body" represents a protective  
15 group which is cleaved by a biological method such as  
hydrolysis in a human body and produces free acids or the  
salts thereof. Whether a derivative is cleavable or not can  
be determined by administering it to an experimental animal  
such as a rat or mouse by an intravenous injection, analyzing  
20 a body fluid of the animal to detect the original compound or  
the pharmacologically acceptable salt thereof.

"The protective group cleavable by a biological method  
such as hydrolysis in a living body" relating to "the ester  
of the hydroxy group" may include a 1-("aliphatic acyl"oxy)  
25 "lower alkyl group" such as formyloxymethyl, acetoxymethyl,  
dimethylaminoacetoxymethyl, propionyloxymethyl,  
butyryloxymethyl, pivaloyloxymethyl, valeryloxymethyl,  
isovaleryloxymethyl, hexanoyloxymethyl, 1-formyloxyethyl, 1-  
acetoxyethyl, 1-propionyloxyethyl, 1-butyryloxyethyl, 1-  
30 pivaloyloxyethyl, 1-valeryloxyethyl, 1-isovaleryloxyethyl, 1-  
hexanoyloxyethyl, 1-formyloxypropyl, 1-acetoxypentyl, 1-  
propionyloxypropyl, 1-butyryloxypropyl, 1-pivaloyloxypropyl,  
1-valeryloxypropyl, 1-isovaleryloxypropyl, 1-  
hexanoyloxypropyl, 1-acetoxybutyl, 1-propionyloxybutyl, 1-  
35 butyryloxybutyl, 1-pivaloyloxybutyl, 1-acetoxypentyl, 1-  
propionyloxypropyl, 1-butyryloxypropyl, 1-pivaloyloxypropyl,  
1-pivaloyloxyhexyl and etc.; a 1-("aliphatic acyl"thio)

"lower alkyl group" such as formylthiomethyl,  
 acetylthiomethyl, dimethylaminoacetylthiomethyl,  
 propionylthiomethyl, butyrylthiomethyl, pivaloylthiomethyl,  
 valerylthiomethyl, isovalerylthiomethyl, hexanoylthiomethyl,  
 5 1-formylthioethyl, 1-acetylthioethyl, 1-propionylthioethyl,  
 1-butyrylthioethyl, 1-pivaloylthioethyl, 1-valerylthioethyl,  
 1-isovalerylthioethyl, 1-hexanoylthioethyl, 1-  
 formylthiopropyl, 1-acetylthiopropyl, 1-propionylthiopropyl,  
 1-butyrylthiopropyl, 1-pivaloylthiopropyl, 1-  
 10 valerylthiopropyl, 1-isovalerylthiopropyl, 1-  
 hexanoylthiopropyl, 1-acetylthiobutyl, 1-propionylthiobutyl,  
 1-butyrylthiobutyl, 1-pivaloylthiobutyl, 1-acetylthiopentyl,  
 1-propionylthiopentyl, 1-butyrylthiopentyl, 1-  
 pivaloylthiopentyl, 1-pivaloylthiohexyl and etc.; a 1-  
 15 (acyloxy) "lower alkyl group" such as a 1-  
 ("cycloalkyl"carbonyloxy) "lower alkyl group", e.g.,  
 cyclopentylcarbonyloxymethyl, cyclohexylcarbonyloxymethyl, 1-  
 cyclopentylcarbonyloxyethyl, 1-cyclohexylcarbonyloxyethyl, 1-  
 cyclopentylcarbonyloxypropyl, 1-cyclohexylcarbonyloxypropyl,  
 20 1-cyclopentylcarbonyloxybutyl and 1-  
 cyclohexylcarbonyloxybutyl, 1-("aromatic acyl"oxy) "lower  
 alkyl group", e.g., benzoyloxymethyl;  
 (alkoxycarbonyloxy)alkyl group such as  
 methoxycarbonyloxymethyl, ethoxycarbonyloxymethyl,  
 25 propoxycarbonyloxymethyl, isopropoxycarbonyloxymethyl,  
 butoxycarbonyloxymethyl, isobutoxycarbonyloxymethyl,  
 pentyloxycarbonyloxymethyl, hexyloxycarbonyloxymethyl,  
 cyclohexyloxycarbonyloxymethyl,  
 cyclohexyloxycarbonyloxy(cyclohexyl)methyl, 1-  
 30 (methoxycarbonyloxy)ethyl, 1-(ethoxycarbonyloxy)ethyl, 1-  
 (propoxycarbonyloxy)ethyl, 1-(isopropoxycarbonyloxy)ethyl, 1-  
 (butoxycarbonyloxy)ethyl, 1-(isobutoxycarbonyloxy)ethyl, 1-  
 (tert-butoxycarbonyloxy)ethyl, 1-(pentyloxycarbonyloxy)ethyl,  
 1-(hexyloxycarbonyloxy)ethyl, 1-  
 35 (cyclopentyloxycarbonyloxy)ethyl, 1-  
 (cyclopentyloxycarbonyloxy)propyl, 1-  
 (cyclohexyloxycarbonyloxy)propyl, 1-

(cyclopentyloxy)carbonyloxy)butyl, 1-  
(cyclohexyloxy)carbonyloxy)butyl, 1-  
(cyclohexyloxy)carbonyloxy)ethyl, 1-(ethoxycarbonyloxy)propyl,  
2-(methoxycarbonyloxy)ethyl, 2-(ethoxycarbonyloxy)ethyl, 2-  
5 (propoxycarbonyloxy)ethyl, 2-(isopropoxycarbonyloxy)ethyl, 2-  
(butoxycarbonyloxy)ethyl, 2-(isobutoxycarbonyloxy)ethyl, 2-  
(pentyloxy)carbonyloxy)ethyl, 2-(hexyloxy)carbonyloxy)ethyl, 1-  
(methoxycarbonyloxy)propyl, 2-(ethoxycarbonyloxy)propyl, 1-  
(propoxycarbonyloxy)propyl, 1-(isopropoxycarbonyloxy)propyl,  
10 1-(butoxycarbonyloxy)propyl, 1-(isobutoxycarbonyloxy)propyl,  
1-(pentyloxy)carbonyloxy)propyl, 1-(hexyloxy)carbonyloxy)propyl,  
1-(methoxycarbonyloxy)butyl, 1-(ethoxycarbonyloxy)butyl, 1-  
(propoxycarbonyloxy)butyl, 1-(isopropoxycarbonyloxy)butyl, 1-  
(butoxycarbonyloxy)butyl, 1-(isobutoxycarbonyloxy)butyl, 1-  
15 (methoxycarbonyloxy)pentyl, 1-(ethoxycarbonyloxy)pentyl, 1-  
(methoxycarbonyloxy)hexyl, 1-(ethoxycarbonyloxy)hexyl and  
etc.; "a phthalidyl group" such as phthalidyl,  
dimethylphthalidyl and dimethoxyphthalidyl; "a  
carbonyloxyalkyl group" such as an oxodioxolenylmethyl group,  
20 e.g., (5-phenyl-2-oxo-1,3-dioxolen-4-yl)methyl, [5-(4-  
methylphenyl)-2-oxo-1,3-dioxolen-4-yl]methyl, [5-(4-  
methoxyphenyl)-2-oxo-1,3-dioxolen-4-yl]methyl, [5-(4-  
fluorophenyl)-2-oxo-1,3-dioxolen-4-yl]methyl, [5-(4-  
chlorophenyl)-2-oxo-1,3-dioxolen-4-yl]methyl, (2-oxo-1,3-  
25 dioxolen-4-yl)methyl, (5-methyl-2-oxo-1,3-dioxolen-4-  
yl)methyl, (5-ethyl-2-oxo-1,3-dioxolen-4-yl)methyl, (5-  
propyl-2-oxo-1,3-dioxolen-4-yl)methyl, (5-isopropyl-2-oxo-  
1,3-dioxolen-4-yl)methyl, (5-butyl-2-oxo-1,3-dioxolen-4-  
yl)methyl and etc.; the above-mentioned "aliphatic acyl  
30 group"; the above-mentioned "aromatic acyl group"; "a half  
ester salt residual group of succinic acid"; "a phosphoric  
acid ester salt residual group"; "ester-forming residual  
group of amino acid, and etc."; a carbamoyl group; a  
carbamoyl group substituted by 1 or 2 lower alkyl groups; a  
35 carboxyl "lower alkyl group" dithioethyl group such as 2-  
carboxyl ethyldithioethyl, 3-carboxyl propyldithioethyl, 4-  
carboxyl butyldithioethyl, 5-carboxyl pentyldithioethyl, 6-

carboxyl hexyldithioethyl and etc.; and "a lower alkyl group" dithioethyl group such as methyldithioethyl, ethyldithioethyl, propyldithioethyl, butyldithioethyl, pentyldithioethyl, hexyldithioethyl and etc.

5       Whereas, "the protective group cleavable by a biological method such as hydrolysis in a living body" relating to the "ester of the carboxy group" may specifically include "alkoxy lower alkyl group" such as a lower alkoxy lower alkyl group, e.g., methoxymethyl, 1-ethoxyethyl, 1-methyl-1-methoxyethyl, 10 1-(isopropoxy)ethyl, 2-methoxyethyl, 2-ethoxyethyl, 1,1-dimethyl-1-methoxymethyl, ethoxymethyl, n-propoxymethyl, isopropoxymethyl, n-butoxymethyl, tert-butoxymethyl and etc., a lower alkoxyated lower alkoxy lower alkyl group, e.g., 2-methoxyethoxymethyl, an "aryl"oxy "lower alkyl group", e.g., 15 phenoxymethyl and a halogenated lower alkoxy lower alkyl group, e.g., 2,2,2-trichloroethoxymethyl, bis(2-chloroethoxy)methyl and etc.; a "lower alkoxy" carbonyl "lower alkyl group" such as methoxycarbonylmethyl; a cyano "lower alkyl group" such as cyanomethyl and 2-cyanoethyl; a 20 "lower alkyl" thiomethyl group such as methylthiomethyl and ethylthiomethyl; an "aryl" thiomethyl group such as phenylthiomethyl, naphthylthiomethyl and etc.; a "lower alkyl" sulfonyl "lower alkyl group" which may be substituted by halogen atoms such as 2-methanesulfonylethyl and 2- 25 trifluoromethanesulfonylethyl; an "aryl" sulfonyl "lower alkyl group" such as 2-benzenesulfonylethyl and 2-toluenesulfonylethyl; an acyloxy "lower alkyl group" such as an "aliphatic acyl" oxy "lower alkyl group", e.g., formyloxymethyl, acetoxymethyl, propionyloxymethyl, 30 butyryloxymethyl, pivaloyloxymethyl, valeryloxymethyl, isovaleryloxymethyl, hexanoyloxymethyl, 1-formyloxyethyl, 1-acetoxyethyl, 1-propionyloxyethyl, 1-butyryloxyethyl, 1-pivaloyloxyethyl, 1-valeryloxyethyl, 1-isovaleryloxyethyl, 1-hexanoyloxyethyl, 2-formyloxyethyl, 2-acetoxyethyl, 2- 35 propionyloxyethyl, 2-butyryloxyethyl, 2-pivaloyloxyethyl, 2-valeryloxyethyl, 2-isovaleryloxyethyl, 2-hexanoyloxyethyl, 1-formyloxypropyl, 1-acetoxypropyl, 1-propionyloxypropyl, 1-

butyryloxypropyl, 1-pivaloyloxypropyl, 1-valeryloxypropyl, 1-  
 isovaleryloxypropyl, 1-hexanoyloxypropyl, 1-acetoxybutyl, 1-  
 propionyloxybutyl, 1-butyryloxybutyl, 1-pivaloyloxybutyl, 1-  
 acetoxypentyl, 1-propionyloxypentyl, 1-butyryloxypentyl, 1-  
 5 pivaloyloxypentyl, 1-pivaloyloxyhexyl and etc., a  
 "cycloalkyl" carbonyloxy "lower alkyl group", e.g.,  
 cyclopentylcarbonyloxymethyl, cyclohexylcarbonyloxymethyl, 1-  
 cyclopentylcarbonyloxyethyl, 1-cyclohexylcarbonyloxyethyl, 1-  
 cyclopentylcarbonyloxypropyl, 1-cyclohexylcarbonyloxypropyl,  
 10 1-cyclopentylcarbonyloxybutyl, 1-cyclohexylcarbonyloxybutyl  
 and etc., and an "aromatic acyl" oxy "lower alkyl group",  
 e.g., benzoyloxymethyl; an (alkoxycarbonyloxy)alkyl group  
 such as methoxycarbonyloxymethyl, ethoxycarbonyloxymethyl,  
 propoxycarbonyloxymethyl, isopropoxycarbonyloxymethyl,  
 15 butoxycarbonyloxymethyl, isobutoxycarbonyloxymethyl,  
 pentyloxycarbonyloxymethyl, hexyloxycarbonyloxymethyl,  
 cyclohexyloxycarbonyloxymethyl,  
 cyclohexyloxycarbonyloxy(cyclohexyl)methyl, 1-  
 (methoxycarbonyloxy)ethyl, 1-(ethoxycarbonyloxy)ethyl, 1-  
 20 (propoxycarbonyloxy)ethyl, 1-(isopropoxycarbonyloxy)ethyl, 1-  
 (butoxycarbonyloxy)ethyl, 1-(isobutoxycarbonyloxy)ethyl, 1-  
 (tert-butoxycarbonyloxy)ethyl, 1-(pentyloxycarbonyloxy)ethyl,  
 1-(hexyloxycarbonyloxy)ethyl, 1-  
 (cyclopentyloxycarbonyloxy)ethyl, 1-  
 25 (cyclopentyloxycarbonyloxy)propyl, 1-  
 (cyclohexyloxycarbonyloxy)propyl, 1-  
 (cyclopentyloxycarbonyloxy)butyl, 1-  
 (cyclohexyloxycarbonyloxy)butyl, 1-  
 (cyclohexyloxycarbonyloxy)ethyl, 1-(ethoxycarbonyloxy)propyl,  
 30 2-(methoxycarbonyloxy)ethyl, 2-(ethoxycarbonyloxy)ethyl, 2-  
 (propoxycarbonyloxy)ethyl, 2-(isopropoxycarbonyloxy)ethyl, 2-  
 (butoxycarbonyloxy)ethyl, 2-(isobutoxycarbonyloxy)ethyl, 2-  
 (pentyloxycarbonyloxy)ethyl, 2-(hexyloxycarbonyloxy)ethyl, 1-  
 (methoxycarbonyloxy)propyl, 1-(ethoxycarbonyloxy)propyl, 1-  
 35 (propoxycarbonyloxy)propyl, 1-(isopropoxycarbonyloxy)propyl,  
 1-(butoxycarbonyloxy)propyl, 1-(isobutoxycarbonyloxy)propyl,  
 1-(pentyloxycarbonyloxy)propyl, 1-(hexyloxycarbonyloxy)propyl,

1-(methoxycarbonyloxy)butyl, 1-(ethoxycarbonyloxy)butyl, 1-(propoxycarbonyloxy)butyl, 1-(isopropoxycarbonyloxy)butyl, 1-(butoxycarbonyloxy)butyl, 1-(isobutoxycarbonyloxy)butyl, 1-(methoxycarbonyloxy)pentyl, 1-(ethoxycarbonyloxy)pentyl, 1-(methoxycarbonyloxy)hexyl, 1-(ethoxycarbonyloxy)hexyl and etc.; "a carbonyloxyalkyl group" such as an oxodioxolenylmethyl group, e.g., (5-phenyl-2-oxo-1,3-dioxolen-4-yl)methyl, [5-(4-methylphenyl)-2-oxo-1,3-dioxolen-4-yl]methyl, [5-(4-methoxyphenyl)-2-oxo-1,3-dioxolen-4-yl]methyl, [5-(4-fluorophenyl)-2-oxo-1,3-dioxolen-4-yl]methyl, [5-(4-chlorophenyl)-2-oxo-1,3-dioxolen-4-yl]methyl, (2-oxo-1,3-dioxolen-4-yl)methyl, (5-methyl-2-oxo-1,3-dioxolen-4-yl)methyl, (5-ethyl-2-oxo-1,3-dioxolen-4-yl)methyl, (5-propyl-2-oxo-1,3-dioxolen-4-yl)methyl, (5-isopropyl-2-oxo-1,3-dioxolen-4-yl)methyl, (5-butyl-2-oxo-1,3-dioxolen-4-yl)methyl and etc.; "a phthalidyl group" such as phthalidyl, dimethylphthalidyl and dimethoxyphthalidyl; "an aryl group" such as phenyl, indanyl and etc.; the above-mentioned "lower alkyl group"; and a straight or branched alkylthio group having 1 to 6 carbon atoms such as methylthio, ethylthio, n-propylthio, isopropylthio, n-butylthio, isobutylthio, s-butylthio, tert-butylthio, n-pentylthio, isopentylthio, 2-methylbutylthio, neopentylthio, 1-ethylpropylthio, n-hexylthio, isohexylthio, 4-methylpentylthio, 3-methylpentylthio, 2-methylpentylthio, 1-methylpentylthio, 3,3-dimethylbutylthio, 2,2-dimethylbutylthio, 1,1-dimethylbutylthio, 1,2-dimethylbutylthio, 1,3-dimethylbutylthio, 2,3-dimethylbutylthio, 2-ethylbutylthio and etc., preferably "an alkylthio group" having 1 to 4 carbon atoms; a "carboxyalkyl group" such as carboxymethyl; and an "amide-forming residual group of an amino acid" such as phenylalanine.

"Other derivatives" means an ether derivative or a carbamoyloxy derivative in the case where Compound (I) of the present invention has "a hydroxy group", or means an amide derivative in the case where the Compound (I) of the present invention has "an amino group", that are decomposed in the

living body to form their respective original "hydroxy group" or "amino group". A determination whether or not a derivative is such a kind, can be made by administering it by intravenous injection to an experimental animal such as a rat or a mouse, analyzing the animal's body fluids afterward, to detect the original compound or its pharmacologically acceptable salt.

Compound (I) of the present invention may occasionally have an asymmetrical carbon in the molecule, and stereoisomers of R and S-configuration sometimes exist. Each of the stereoisomers or mixtures containing such stereoisomers in an arbitrary proportion thereof are all included in the present invention.

Moreover, the external preparation of the present invention is an external preparation which comprises at least one compound of the above nitroimidazole derivatives and at least one agent selected from the group consisting of an antimycotic agent, antibacterial agent, sulfa, immunosuppressant, anti-inflammatory drug, antibiotic, antiviral agent, metabolic antagonist, antihistamine, tissue repair promoter, vitamin, antiallergic, local anesthetic, hair agent and steroid being administered simultaneously or separately with an interval.

In the above, there are no particular restrictions on "administered simultaneously" provided it is an administration form that can be administered at nearly the same time, it is preferable to administer the preparation in the form of a single composition.

In the above, there are no particular restrictions on "administered separately with an interval" provided it is an administration form that can be administered separately with an interval. It refers to, for example, administration of a nitroimidazole derivative on day 1 followed by administration on day 2 of a preparation containing at least one agent selected from the group consisting of an antimycotic agent, antibacterial agent, sulfa, immunosuppressant, anti-inflammatory agent, antibiotic, antiviral agent, metabolic

antagonist, antihistamine, tissue repair promoter, vitamin, antiallergic, local anesthetic, hair agent and steroid, or to initially administering a nitroimidazole derivative and then with a predetermined interval, administering a preparation  
5 containing at least one agent selected from the group consisting of an antimycotic agent, antibacterial agent, sulfa, immunosuppressant, anti-inflammatory agent, antibiotic, antiviral agent, metabolic antagonist, antihistamine, tissue repair promoter, vitamin, antiallergic, local anesthetic,  
10 hair agent, steroid and the like.

Among at least one agent selected from an antimycotic agent, antibacterial agent, sulfa, immunosuppressant, anti-inflammatory agent, antibiotic, antiviral agent, metabolic  
15 antagonist, antihistamine, tissue repair promoter, vitamin, antiallergic, local anesthetic, hair agent and steroid in the above description, an antimycotic agent, immunosuppressant, steroid and their combinations are preferable for an external preparation for the therapeutic or prophylactic treatment of atopic dermatitis, while an immunosuppressant, steroid and  
20 the combination of antimycotic agent and steroid are more preferable. An antimycotic agent, immunosuppressant, vitamin, antiallergic, steroid and their combinations are preferable for an external preparation for the therapeutic or prophylactic treatment of psoriasis, while an  
25 immunosuppressant, vitamin and steroid are more preferable. An antibacterial agent is preferable for an external preparation for the therapeutic or prophylactic treatment of tinea, while an antibiotic is preferable for an external preparation for the therapeutic or prophylactic treatment of  
30 suppurative skin diseases.

In the above, the agent of at least one agent selected from the group consisting of an antimycotic agent, antibacterial agent, sulfa, immunosuppressant, anti-inflammatory agent, antibiotic, antiviral agent, metabolic  
35 antagonist, antihistamine, tissue repair promoter, vitamin, antiallergic, local anesthetic, hair agent and steroid, is preferably used in a concentration at which the agent itself

does not demonstrate any pharmacological effect.

Determination on whether or not a concentration is at a level at which a pharmacological effect is demonstrated can be easily done by a person with ordinary skill in the art using commonly known means (such as comparative studies in humans or animals).

There are no particular restrictions on the content in a preparation in the case of containing in an external preparation an antimycotic agent, antibacterial agent, sulfa, immunosuppressant, anti-inflammatory agent, antibiotic, antiviral agent, metabolic antagonist, antihistamine, tissue repair promoter, vitamin, antiallergic, local anesthetic, hair agent or steroid, provided that the concentration is not that at which adverse side effects are exhibited. Based on the weight of the preparation, the content is preferably 0.0005 to 2 wt%, and more preferably 0.01 to 0.5 wt% for an antimycotic agent, preferably 0.001 to 5 wt%, and more preferably 0.01 to 0.5 wt% for an antibacterial agent, preferably 0.001 to 5 wt%, and more preferably 0.01 to 0.5 wt% for a sulfa, preferably 0.001 to 5 wt%, and more preferably 0.01 to 0.1 wt% for an immunosuppressant, preferably 0.001 to 5 wt%, and more preferably 0.005 to 0.5 wt% for an anti-inflammatory agent, preferably 0.0001 to 5 wt%, and more preferably 0.001 to 0.1 wt% for an antibiotic, preferably 0.01 to 5 wt%, and more preferably 0.1 to 1 wt% for an antiviral agent, preferably 0.01 to 5 wt%, and more preferably 0.01 to 0.5 wt% for a metabolic antagonist, preferably 0.001 to 10 wt%, and more preferably 0.01 to 5 wt% for an antihistamine, preferably 0.1 to 20 wt%, and more preferably 0.1 to 5 wt% for a tissue repair promoter, preferably 0.000001 to 0.005 wt%, and more preferably 0.00001 to 0.001 wt% for a vitamin, preferably 0.001 to 5 wt%, and more preferably 0.01 to 0.5 wt% for an antiallergic, preferably 0.001 to 5 wt%, and more preferably 0.01 to 1 wt% for a local anesthetic, preferably 0.01 to 10 wt%, and more preferably 0.1 to 2 wt% for a hair agent, and preferably 0.001 to 1 wt%, and more preferably 0.001 to 0.1 wt% for a

steroid.

There are no particular restrictions on the above antimycotic agent provided it is an agent that is used for the treatment of pathogenic molds and deep mycoses, examples of which include imidazole compounds such as croconazole hydrochloride, neticonazole hydrochloride, clotrimazole, ketoconazole, isoconazole nitrate, econazole nitrate, oxiconazole nitrate, sulconazole nitrate, miconazole nitrate, thioconazole, bifonazole and lanoconazole, as well as amorolfine hydrochloride, terbinafine hydrochloride, butenafine hydrochloride, ciclopirox olamine, tolciolate, tolnaftate and the like.

There are no particular restrictions on the above antibacterial agent provided it is an agent that has efficacy against pathogenic microorganisms (including Gram positive cocci and bacilli, and Gram negative cocci and bacilli), examples of which include enoxacin, methyl rosaniline chloride, ciprofloxacin hydrochloride, lomefloxacin hydrochloride, ofloxacin, cinoxacin, sparfloxacin, tosufloxacin tosilate, nalidixic acid, norfloxacin, pipemidic acid trihydrate, piromidic acid, fleroxacin, levofloxacin, etc. and their derivatives.

There are no particular restrictions on the above sulfa provided it is used routinely, examples of which include acetylsulfamethoxazole, salazosulfapyridine, sulfadiazine, sulfadiazine silver, sulfadimethoxine, sulfathiazole, sulfaphenazole, sulfamethoxazole, sulfamethoxypyridazine, sulfamethopyradine, sulfamethomidine, sulfamethizole, sulfameradine, sulfamonomethoxine, sulfisoxazole, sulfisomidin, sulfisomidin sodium, homosulfamine, etc. and their derivatives.

There are no particular restrictions on the above immunosuppressant provided that an agent suppresses immune rejection reactions, examples of which include pimecrolimus, sirolimus, eberolimus, cyclosporin, tacrolimus, glibelimus hydrochloride, mizoribine, FTY-720 (2-amino-2-(2-(4-octylphenyl)ethyl)propane-1,3-diol hydrochloride) and etc.

There are no particular restrictions on the above anti-inflammatory agent provided it is used routinely, examples of which include actarit, azulene, acemetacin, aspirin,

alclofenac, alminoprofen, amfenac sodium, ampiroxicam,

5 ibuprofen, ibuprofenpiconol, indometacin, indometacin

farnesil, ufenamate, etodolac, epirizol, emorfazone,

tiaramide hydrochloride, tinoridine hydrochloride,

buprenorphine hydrochloride, pentazocine hydrochloride,

enfenam, oxaprozin, glycyrrhetic acid, crotamiton, ketoprofen,

10 zaltoprofen, diflunisal, diclofenac sodium, suprofen,

sulindac, tiaprofen, tenoxicam, trimethine sodium, nabumeton,

naproxen, nifurmic acid, piroxicam, phenacetin,

phenylbutazone, phenoprofen calcium, felbinac, fenbufen,

bucolome, bufexamac, pranoprofen, flurbiprofen, floctafenine,

15 dimethothiazine mesilate, methiazine, bendazac, heparin

analogues, proglumetacin maleate, meclofenam, mefenamic acid,

loxoprofen sodium, lobenzarit disodium, vaccinia virus

inoculated rabbit inflammatory skin extract, etc. and

derivatives thereof.

20 The above antibiotic means a substance that inhibits the growth of microorganisms, examples of which include

acetylkitasamycin, acetylspiramycin, amphotericin B,

amoxicillin, ampicillin, kanamycin monosulfate, erythromycin

ethyl succinate, erythromycin, erythromycin estorate,

25 aclarubicin hydrochloride, oxytetracycline hydrochloride,

clindamycin hydrochloride, cefetamet pivoxil hydrochloride,

cefotiam hexetil hydrochloride, cefcapene pivoxil

hydrochloride, cefmenoxime hydrochloride, talampicillin

hydrochloride, tetracycline hydrochloride,

30 demethylchlortetracycline hydrochloride, tetracycline

hydrochloride, vancomycin hydrochloride, doxycycline

hydrochloride, doxorubicin hydrochloride, bacampicillin

hydrochloride, clindamycin palmitate hydrochloride,

vancomycin hydrochloride, pivmecillinam hydrochloride,

35 bleomycin hydrochloride, minocycline hydrochloride,

lincomycin hydrochloride, lenampicillin hydrochloride,

carbenicillin sodium, kitasamycin, potassium clavulanate,

clarithromycin, griseofulvin, cloxaciline sodium,  
 chloramphenicol, colistin sodium methanesulfonate,  
 cycloserine, midecamycin acetate, ciclacillin, dicloxacillin  
 sodium, siccanin, josamycin, erythromycin stearate,  
 5 sulbenicillin sodium, cefaclor, cefazolin, propylene glycol  
 cefatrizine, cefadroxil, cefapirin, cefamandole sodium,  
 cefalexin, cefalotin sodium, cefaloridine, cefixime,  
 ceftizoxime sodium, cefotaxime sodium, cefotetan, cefoperazone  
 sodium, cefditoren pivoxil, cefdinir, cefsulodin sodium,  
 10 ceftizoxime sodium, ceftibuten, cefteteram pivoxil, cefpiramide  
 sodium, cefbuperazone sodium, cefpodoxime proxetil,  
 cefmetazole sodium, cefradine, cefroxadine, cefuroxime axetil,  
 cefuroxime sodium, ticarcillin sodium, tetracycline,  
 sultamicillin tosilate, tobramycin, trichomycin, nystatin,  
 15 variotin, chloramphenicol palmitate, piperacillin sodium,  
 pimarcin, faropenem sodium, josamycin propionate,  
 phenethicillin potassium, phenoxymethylpenicillin potassium,  
 benzylpenicillin potassium, benzylpenicillin benzathine,  
 fosfomicin calcium, mitomycin C, midecamycin, tetracycline  
 20 metaphosphate, latamoxef sodium, rifampicin, astromicin  
 sulfate, amikacin sulfate, kanamycin sulfate, gentamicin  
 sulfate, sisomicin sulfate, dibekacin sulfate, streptomycin  
 sulfate, netilmicin sulfate, fradiomycin sulfate, bleomycin  
 sulfate, bekanamycin sulfate, peplomycin sulfate, polymyxin B  
 25 sulfate, micronomicin sulfate, ribostamycin sulfate,  
 clindamycin phosphate, roxithromycin, rokitamycin, etc. and  
 derivatives thereof.

The above antiviral agent means an agent that is  
 specific for viruses, examples of which include aciclovir,  
 30 ganciclovir, sanilvudine, zalcitabine, didanosine, zidovudine,  
 nevirapine, saquinavir mesilate, nelfinavir mesilate,  
 lamivudine, ritonavir, indinavir sulfate, etc. and the  
 addition and substitution products of their salts.

There are no particular restrictions on the above  
 35 metabolic antagonist provided it is used routinely, examples  
 of which include actinomycin D, L-asparaginase, aceglatone,  
 ubenimex, uracil, etoposide, enocitabine, aclarubicin

hydrochloride, idarubicin hydrochloride, irinotecan  
hydrochloride, epirubicin hydrochloride, dunorubicin  
hydrochloride, doxorubicin hydrochloride, pirarubicin  
hydrochloride, fadrozole hydrochloride hydrate, bleomycin  
5 hydrochloride, procarbazine hydrochloride, mitoxantrone  
hydrochloride, carboplatin, carmofur, tamoxifen citrate,  
toremifene citrate, cyclophosphamide, cisplatin, sizofiran,  
cytarabine, cytarabine ocfosfate, zinostatin stimalamer,  
vinorelbine ditartrate, sobuzoxane, thiotepa, tegafur,  
10 doxifluridine, docetaxel hydrate, tretinoin, neocarzinostatin,  
nedaplatin, paclitaxel, bicalutamide, hydroxycarbamide,  
fosfestrol, busulfan, fluorouracil, flutamide,  
propylthiouracil, pentostatin, porfimer sodium,  
methyltestosterone, mepitiostane, G-mercaptapurine riboside,  
15 mercaptopurine, methotrexate, melphalan, Streptococcus  
extract, peplomycin sulfate, vincristine sulfate, vinblastine  
sulfate, lentinan, etc. and derivatives thereof.

There are no particular restrictions on the above  
antihistamine provided it is an agent that is specifically  
20 antagonistic for histamine, examples of which include  
cyproheptadine hydrochloride, diphenhydramine hydrochloride,  
triprolidine hydrochloride, hydroxidine hydrochloride,  
promethazine hydrochloride, homochlorcyclizine hydrochloride,  
cimetidine, alimemazine tartrate, diphenhydramine tannate,  
25 diphenylpyraline teoclote, hydroxidine pamoate, famotidine,  
chlorpheniramine maleate, clemastine fumarate, mequitazine,  
etc. and derivatives thereof.

There are no particular restrictions on the above tissue  
repair promoter provided it is an agent that promotes tissue  
30 repair, examples of which include extract of calves blood,  
EGF and their derivatives.

The above vitamin refers to vitamins which demonstrate  
vitamin-like action, examples of which include vitamin D3  
analogues such as tacalcitol, mexacalcitol, calcipotriol, and  
35 ferecalcitol, vitamin A analogues such as adaparene,  
tazarotene, alitretinoin, and etretinate, as well as vitamin  
A, vitamin B group, vitamin C, vitamin D and vitamin E.

There are no particular restrictions on the above antiallergic provided it is used routinely, examples of which include astemizole, amlexanox, ibudilast, ebastine, azelastine hydrochloride, epinastine hydrochloride, ozagrel hydrochloride, ceterizine hydrochloride, oxatomide, sodium cromoglicate, seratroast, tazanast, terfenadine, suplastat tosilate, tranilast, emedastine difumarate, ketotifen fumarate, pranlukast hydrate, pemirolast potassium, repirinast, etc. and derivatives thereof.

The above local anesthetic means a drug that is able to anesthetize perception and movement at the location at which it is applied, examples of which include ethyl aminobenzoate, oxybuprocaine hydrochloride, dibucaine hydrochloride, tetracaine hydrochloride, diethylaminoethyl parabutyl-aminobenzoate hydrochloride, procaine hydrochloride, mepivacaine hydrochloride, lidocaine hydrochloride, oxethazaine, lidocaine and etc. and derivatives thereof.

There are no particular restrictions on the above hair agent provided it is used routinely, examples of which include asunaron, carpronium chloride and minoxidil.

There are no particular restrictions on the above steroid provided it is an agent that exhibits action resembling steroid hormones secreted from the adrenal cortex, examples of which include amcinonide, oxymetholone, potassium canrenoate, prednisolone valerate acetate, diflucortolone valerate, dexamethasone valerate, betamethasone valerate, hydrocortisone succinate, prednisolone succinate, chlormadinone acetate, cortisone acetate, diflorasone diacetate, hydrocortisone acetate, paramethasone acetate, fludrocortisone acetate, prednisolone acetate, methenolone acetate, difluprednate, betamethasone dipropionate, dexamethasone, triamcinolone, triamcinolone acetonide, halcinonide, hydrocortisone, flumetasone pivalate, prednisolone farnesylate gel, budesonide, mometasone furoate, fluocinonide, fluocinolone acetonide, fluorometholone, fludroxycortide, prednisolone, alclometasone dipropionate, clobetasol propionate, dexamethasone propionate, deprodone

propionate, beclometasone dipropionate, betamethasone, methylprednisolone, clobetasone butyrate, hydrocortisone butyrate, hydrocortisone butyrate propionate, betamethasone butyrate propionate, hydrocortisone sodium phosphate, 5 betamethasone sodium phosphate, etc. and derivatives thereof.

Preferable examples of the above nitroimidazole derivatives include the above-mentioned (1) through (9), and more preferable examples include metronidazole and tinidazole.

10 In addition, the external preparation for skin diseases of the present invention is preferably an external preparation for skin diseases that contains crotamiton. The containing of crotamiton has the effect of fast-action of antiamyctic effects, increasing solubility of nitroimidazole derivative, and improving stability of the external 15 preparation.

Preferable examples of skin diseases of the external preparation for therapeutic or prophylactic treatment or amelioration of skin diseases of the present invention include:

- 20 (10) atopic dermatitis,  
(11) facial atopic dermatitis,  
(12) pediatric atopic dermatitis,  
(13) skin blotches, pigmentation or scars,  
(14) psoriasis,  
25 (15) hircus, body odor or osmidrosis,  
(16) contact dermatitis, plant dermatitis or insect bites,  
(17) dermal pruritis or drug rash,  
(18) chilblain,  
(19) erythroderma,  
30 (20) tinea,  
(21) suppurative skin diseases,  
(22) bed sores,  
(23) wounds, and  
(24) palmoplantar pustulosis, lichen planus, lichen nitidus,  
35 pityriasis rubra pilaris, pityriasis rosea, erythema  
(including polymorphic exudative erythema, nodular erythema and Darier's erythema annulare centrifugum), chronic discoid

lupus erythematosus, drug rash and toxic rash, alopecia areata, burns (including scars and keloids), pemphigus, Duhring dermatitis herpetiformis (including pemphigoid), seborrheic dermatitis, dermal stomatitis, Candidiasis

5 (including interdigital erosion, intertrigo, dermal Candidiasis, infantile parasitic erythema, perionychia and vaginal Candidiasis) and tinea versicolor, while more preferable examples include (10), (11), (12), (13), (14) and (15). In addition, during the above therapeutic or

10 prophylactic treatment or amelioration of skin diseases, an external preparation is also preferable in which a

nitroimidazole derivative is arbitrarily selected, and at least one compound of the nitroimidazole derivatives and at least one agent selected from the group consisting of an

15 antimycotic agent, antibacterial agent, sulfa, immunosuppressant, anti-inflammatory agent, antibiotic, antiviral agent, metabolic antagonist, antihistamine, tissue repair promoter, vitamin, antiallergic, local anesthetic, hair agent and steroid are administered simultaneously or  
20 separately with an interval, while more preferable examples of external preparations include:

an external preparation for a therapeutic or prophylactic treatment of (10) atopic dermatitis in which the nitroimidazole derivative is metronidazole,

25 an external preparation for a therapeutic or prophylactic treatment of (11) facial atopic dermatitis in which the nitroimidazole derivative is metronidazole,

an external preparation for a therapeutic or prophylactic treatment of (12) pediatric atopic dermatitis in  
30 which the nitroimidazole derivative is metronidazole,

an external preparation for amelioration of (13) skin blotches, pigmentation or scars in which the nitroimidazole derivative is metronidazole,

an external preparation for a therapeutic or  
35 prophylactic treatment of (14) psoriasis in which the nitroimidazole derivative is metronidazole,

an external preparation for a therapeutic or

prophylactic treatment of (15) hircus, body odor or osmidrosis in which the nitroimidazole derivative is metronidazole,

an external preparation for a therapeutic or prophylactic treatment of (10) atopic dermatitis in which the nitroimidazole derivative is metronidazole, and metronidazole and an antimycotic agent, immunosuppressant, steroid or their combination being administered simultaneously or separately with an interval,

an external preparation for a therapeutic or prophylactic treatment of (10) atopic dermatitis in which the nitroimidazole derivative is metronidazole, and metronidazole and immunosuppressant, steroid, or a combination of antimycotic agent and steroid being administered simultaneously or separately with an interval,

an external preparation for a therapeutic or prophylactic treatment of (14) psoriasis in which the nitroimidazole derivative is metronidazole, and metronidazole and an antimycotic agent, immunosuppressant, vitamins, antiallergic, steroid or their combination being administered simultaneously or separately with an interval,

an external preparation for a therapeutic or prophylactic treatment of (14) psoriasis in which the nitroimidazole derivative is metronidazole, and metronidazole and immunosuppressant, vitamins or steroid being administered simultaneously or separately with an interval,

an external preparation for a therapeutic or prophylactic treatment of (10) atopic dermatitis in which the nitroimidazole derivative is tinidazole,

an external preparation for a therapeutic or prophylactic treatment of (11) facial atopic dermatitis in which the nitroimidazole derivative is tinidazole,

an external preparation for a therapeutic or prophylactic treatment of (12) pediatric atopic dermatitis in which the nitroimidazole derivative is tinidazole,

an external preparation for amelioration of (13) skin blotches, pigmentation or scars in which the nitroimidazole

derivative is tinidazole,

an external preparation for a therapeutic or prophylactic treatment of (14) psoriasis in which the nitroimidazole derivative is tinidazole,

5 an external preparation for a therapeutic or prophylactic treatment of (15) hircus, body odor or osmidrosis in which the nitroimidazole derivative is tinidazole,

10 an external preparation for a therapeutic or prophylactic treatment of (10) atopic dermatitis in which the nitroimidazole derivative is tinidazole, and tinidazole and an antimycotic agent, immunosuppressant, steroid or their combination being administered simultaneously or separately with an interval,

15 an external preparation for a therapeutic or prophylactic treatment of (10) atopic dermatitis in which the nitroimidazole derivative is tinidazole, and tinidazole and immunosuppressant, steroid, or a combination of antimycotic agent and steroid being administered simultaneously or  
20 separately with an interval,

An external preparation for a therapeutic or prophylactic treatment of (14) psoriasis in which the nitroimidazole derivative is tinidazole, and tinidazole and an antimycotic agent, immunosuppressant, vitamins,  
25 antiallergic, steroid or their combination being administered simultaneously or separately with an interval, and

An external preparation for a therapeutic or prophylactic treatment of (14) psoriasis in which the  
30 nitroimidazole derivative is tinidazole, and tinidazole and immunosuppressant, vitamins or steroid being administered simultaneously or separately with an interval,

In addition to diseases of humans, diseases of other mammals (such as dogs or cats) are also included in the  
35 targets of the above amelioration, therapeutic or prophylactic treatment.

There are no particular restrictions on the form of the

external preparation for skin diseases of the present invention provided it is used routinely, preferable examples of which include cream, lotion, shampoo, gel, rinse, face lotion, milky lotion, paste, shaving cream, foundation, cologne, pack, ointment, patch, semi-solid, solid or liquid. In the treatment of atopic dermatitis of the head region in particular, an external preparation such as shampoo, gel or rinse is useful since cream or ointment and so forth is difficult to use.

Although there are no particular restrictions on the concentration of the nitroimidazole derivative in the external preparation for skin diseases of the present invention provided it is a concentration at which effects are demonstrated, the concentration is preferably 0.1 to 20 wt%, more preferably 1.0 to 10 wt%, further more preferably 1.5 to 10 wt%, and most preferably 1.5 to 5 wt%, based on the weight of the preparation.

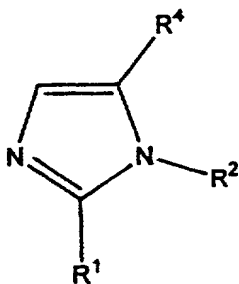
Although there are no particular restrictions on the overall pH of the external preparation for skin diseases of the present invention provided it is pH that is used routinely, pH is preferably 2.0 to 9.0, more preferably 3.0 to 9.0 and particularly preferably 4.0 to 9.0.

In the present invention, a nitroimidazole derivative is used to produce an external preparation for therapeutic or prophylactic treatment of atopic dermatitis, an external preparation for amelioration of skin blotches, pigmentation or scars, an external preparation for therapeutic or prophylactic treatment of psoriasis, and an external preparation for the therapeutic or prophylactic treatment of hircus, body odor or osmidrosis.

In the present invention, therapeutic or prophylactic treatment of atopic dermatitis, amelioration of skin blotches, pigmentation or scars, therapeutic or prophylactic treatment of psoriasis, and therapeutic or prophylactic treatment of hircus, body odor or osmidrosis are performed using an external preparation for skin diseases that contains the nitroimidazole derivative.

Specific compounds included in the nitroimidazole derivatives of the present invention are exemplified in Table 1, but are not limited thereto.

5 In Table 1, Me represents a methyl group, Et an ethyl group, Pr a propyl group, iPr an isopropyl group, Bu a butyl group, Pn a pentyl group, Hx a hexyl group, Ac an acetyl group, Bn a benzyl group, Bz a benzoyl group, Car a carbamoyl group and Mor a morpholino group.



[Table 1]

Compound No.	R <sup>1</sup>	R <sup>4</sup>	R <sup>2</sup>
1	H	NO <sub>2</sub>	Me
2	H	NO <sub>2</sub>	CH <sub>2</sub> OH
3	H	NO <sub>2</sub>	CH <sub>2</sub> OAc
4	H	NO <sub>2</sub>	CH <sub>2</sub> OBn
5	H	NO <sub>2</sub>	CH <sub>2</sub> OBz
6	H	NO <sub>2</sub>	CH <sub>2</sub> SH
7	H	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Me
8	H	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Et
9	H	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
10	H	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
11	H	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
12	H	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
13	H	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
14	H	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Pr
15	H	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> iPr
16	H	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Bu
17	H	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Pn
18	H	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Hx
19	H	NO <sub>2</sub>	CH <sub>2</sub> OCar
20	H	NO <sub>2</sub>	CH <sub>2</sub> NHCO <sub>2</sub> Me
21	H	NO <sub>2</sub>	CH <sub>2</sub> NHC(S)OMe
22	H	NO <sub>2</sub>	Et
23	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OH
24	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OAc
25	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OBn
26	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OBz
27	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SH
28	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
29	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
30	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
31	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
32	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
33	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
34	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH

35	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pr
36	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
37	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Bu
38	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pn
39	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Hx
40	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OCar
41	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> NHCO <sub>2</sub> Me
42	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> NHC(S)OMe
43	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> Mor
44	H	NO <sub>2</sub>	Pr
45	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
46	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
47	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
48	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
49	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
50	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>3</sub>
51	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>3</sub>
52	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>3</sub>
53	H	NO <sub>2</sub>	CH <sub>2</sub> CH(SH)CH <sub>3</sub>
54	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> OMe
55	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> OMe
56	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> OMe
57	H	NO <sub>2</sub>	CH <sub>2</sub> CH(SH)CH <sub>2</sub> OMe
58	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> Cl
59	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> Cl
60	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> Cl
61	H	NO <sub>2</sub>	CH <sub>2</sub> CH(SH)CH <sub>2</sub> Cl
62	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> F
63	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
64	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
65	H	NO <sub>2</sub>	CH <sub>2</sub> CH(CH <sub>3</sub> )SO <sub>2</sub> Et
66	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
67	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
68	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
69	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
70	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
71	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pr

72	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
73	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Bu
74	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pn
75	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Hx
76	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NHCO <sub>2</sub> Me
77	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NHC(S)OMe
78	H	NO <sub>2</sub>	Bu
79	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
80	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
81	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
82	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
83	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
84	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>3</sub>
85	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> CH <sub>3</sub>
86	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> CH <sub>3</sub>
87	H	NO <sub>2</sub>	CH <sub>2</sub> CH(SH)CH <sub>2</sub> CH <sub>3</sub>
88	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> OMe
89	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
90	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
91	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
92	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
93	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
94	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
95	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
96	H	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
97	H	NO <sub>2</sub>	Pn
98	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> OH
99	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> OAc
100	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> OBn
101	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> OBz
102	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SH
103	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
104	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
105	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
106	H	NO <sub>2</sub>	CH <sub>2</sub> CH(SH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
107	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OMe
108	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> Me

109	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> Et
110	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
111	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
112	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
113	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
114	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
115	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> iPr
116	H	NO <sub>2</sub>	Hx
117	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> OH
118	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> OAc
119	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> OBn
120	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> OBz
121	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SH
122	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
123	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
124	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
125	H	NO <sub>2</sub>	CH <sub>2</sub> CH(SH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
126	H	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OMe
127	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> Me
128	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> Et
129	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
130	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
131	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
132	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
133	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
134	H	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> iPr
135	Me	NO <sub>2</sub>	Me
136	Me	NO <sub>2</sub>	CH <sub>2</sub> OH
137	Me	NO <sub>2</sub>	CH <sub>2</sub> OAc
138	Me	NO <sub>2</sub>	CH <sub>2</sub> OBn
139	Me	NO <sub>2</sub>	CH <sub>2</sub> OBz
140	Me	NO <sub>2</sub>	CH <sub>2</sub> SH
141	Me	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Me
142	Me	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Et
143	Me	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
144	Me	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
145	Me	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz

146	Me	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
147	Me	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
148	Me	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Pr
149	Me	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> iPr
150	Me	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Bu
151	Me	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Pn
152	Me	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Hx
153	Me	NO <sub>2</sub>	CH <sub>2</sub> OCar
154	Me	NO <sub>2</sub>	CH <sub>2</sub> NHCO <sub>2</sub> Me
155	Me	NO <sub>2</sub>	CH <sub>2</sub> NHC (S) OMe
156	Me	NO <sub>2</sub>	Et
157	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OH
158	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OAc
159	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OBn
160	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OBz
161	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SH
162	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
163	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
164	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
165	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
166	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
167	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
168	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
169	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pr
170	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
171	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Bu
172	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pn
173	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Hx
174	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OCar
175	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> NHCO <sub>2</sub> Me
176	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> NHC (S) OMe
177	Me	NO <sub>2</sub>	Pr
178	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
179	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
180	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
181	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
182	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH

183	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>3</sub>
184	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>3</sub>
185	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>3</sub>
186	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(SH)CH <sub>3</sub>
187	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> OMe
188	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> OMe
189	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> OMe
190	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(SH)CH <sub>2</sub> OMe
191	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> Cl
192	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> Cl
193	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> Cl
194	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(SH)CH <sub>2</sub> Cl
195	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> F
196	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
197	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
198	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(CH <sub>3</sub> )SO <sub>2</sub> Et
199	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
200	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
201	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
202	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
203	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
204	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pr
205	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
206	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Bu
207	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> PnH <sub>3</sub>
208	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Hx
209	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NHCO <sub>2</sub> Me
210	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NHC(S)OMe
211	Me	NO <sub>2</sub>	Bu
212	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
213	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
214	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
215	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
216	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
217	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>3</sub>
218	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> CH <sub>3</sub>
219	Me	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> CH <sub>3</sub>

220	Me	NO <sub>2</sub>	CH <sub>2</sub> CH (SH) CH <sub>2</sub> CH <sub>3</sub>
221	Me	NO <sub>2</sub>	CH <sub>2</sub> CH (OH) CH <sub>2</sub> CH <sub>2</sub> OMe
222	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
223	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
224	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
225	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
226	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
227	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
228	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
229	Me	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
230	Me	NO <sub>2</sub>	Pn
231	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> OH
232	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> OAc
233	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> OBn
234	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> OBz
235	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SH
236	Me	NO <sub>2</sub>	CH <sub>2</sub> CH (OH) CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
237	Me	NO <sub>2</sub>	CH <sub>2</sub> CH (OAc) CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
238	Me	NO <sub>2</sub>	CH <sub>2</sub> CH (OBz) CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
239	Me	NO <sub>2</sub>	CH <sub>2</sub> CH (SH) CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
240	Me	NO <sub>2</sub>	CH <sub>2</sub> CH (OH) CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OMe
241	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> Me
242	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> Et
243	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
244	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
245	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
246	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
247	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
248	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> iPr
249	Me	NO <sub>2</sub>	Hx
250	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> OH
251	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> OAc
252	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> OBn
253	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> OBz
254	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SH
255	Me	NO <sub>2</sub>	CH <sub>2</sub> CH (OH) CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
256	Me	NO <sub>2</sub>	CH <sub>2</sub> CH (OAc) CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>

257	Me	NO <sub>2</sub>	CH <sub>2</sub> CH (OBz) CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
258	Me	NO <sub>2</sub>	CH <sub>2</sub> CH (SH) CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
259	Me	NO <sub>2</sub>	CH <sub>2</sub> CH (OH) CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OMe
260	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> Me
261	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> Et
262	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
263	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
264	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
265	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
266	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
267	Me	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> iPr
268	Et	NO <sub>2</sub>	Me
269	Et	NO <sub>2</sub>	CH <sub>2</sub> OH
270	Et	NO <sub>2</sub>	CH <sub>2</sub> OAc
271	Et	NO <sub>2</sub>	CH <sub>2</sub> OBn
272	Et	NO <sub>2</sub>	CH <sub>2</sub> OBz
273	Et	NO <sub>2</sub>	CH <sub>2</sub> SH
274	Et	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Me
275	Et	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Et
276	Et	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
277	Et	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
278	Et	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
279	Et	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
280	Et	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
281	Et	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Pr
282	Et	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> iPr
283	Et	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Bu
284	Et	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Pn
285	Et	NO <sub>2</sub>	CH <sub>2</sub> SO <sub>2</sub> Hx
286	Et	NO <sub>2</sub>	CH <sub>2</sub> OCar
287	Et	NO <sub>2</sub>	CH <sub>2</sub> NHCO <sub>2</sub> Me
288	Et	NO <sub>2</sub>	CH <sub>2</sub> NHC (S) OMe
289	Et	NO <sub>2</sub>	Et
290	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OH
291	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OAc
292	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OBn
293	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OBz

294	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SH
295	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
296	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
297	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
298	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
299	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
300	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
301	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
302	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pr
303	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
304	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Bu
305	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pn
306	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Hx
307	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> OCar
308	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> NHCO <sub>2</sub> Me
309	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> NHC (S) OMe
310	Et	NO <sub>2</sub>	Pr
311	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
312	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
313	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
314	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
315	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
316	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (OH) CH <sub>3</sub>
317	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (OAc) CH <sub>3</sub>
318	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (OBz) CH <sub>3</sub>
319	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (SH) CH <sub>3</sub>
320	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (OH) CH <sub>2</sub> OMe
321	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (OAc) CH <sub>2</sub> OMe
322	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (OBz) CH <sub>2</sub> OMe
323	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (SH) CH <sub>2</sub> OMe
324	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (OH) CH <sub>2</sub> Cl
325	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (OAc) CH <sub>2</sub> Cl
326	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (OBz) CH <sub>2</sub> Cl
327	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (SH) CH <sub>2</sub> Cl
328	Et	NO <sub>2</sub>	CH <sub>2</sub> CH (OH) CH <sub>2</sub> F
329	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
330	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et

331	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(CH <sub>3</sub> )SO <sub>2</sub> Et
332	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
333	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
334	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
335	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
336	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
337	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pr
338	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
339	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Bu
340	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pn
341	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Hx
342	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NHCO <sub>2</sub> Me
343	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NHC(S)OMe
344	Et	NO <sub>2</sub>	Bu
345	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
346	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
347	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
348	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
349	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
350	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>3</sub>
351	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> CH <sub>3</sub>
352	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> CH <sub>3</sub>
353	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(SH)CH <sub>2</sub> CH <sub>3</sub>
354	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> OMe
355	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
356	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
357	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
358	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
359	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
360	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
361	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
362	Et	NO <sub>2</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
363	Et	NO <sub>2</sub>	Pn
364	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> OH
365	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> OAc
366	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> OBn
367	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> OBz

368	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SH
369	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
370	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
371	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
372	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(SH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
373	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OMe
374	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> Me
375	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> Et
376	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
377	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
378	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
379	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
380	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
381	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> iPr
382	Et	NO <sub>2</sub>	Hx
383	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> OH
384	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> OAc
385	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> OBn
386	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> OBz
387	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SH
388	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
389	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
390	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
391	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(SH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
392	Et	NO <sub>2</sub>	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OMe
393	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> Me
394	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> Et
395	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
396	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
397	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
398	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
399	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
400	Et	NO <sub>2</sub>	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> iPr
401	CH <sub>2</sub> OCar	NO <sub>2</sub>	H
402	CH <sub>2</sub> OCar	NO <sub>2</sub>	Me
403	CH <sub>2</sub> OCar	NO <sub>2</sub>	Et
404	CH <sub>2</sub> CH <sub>2</sub> OCar	NO <sub>2</sub>	H

405	CH <sub>2</sub> CH <sub>2</sub> OCar	NO <sub>2</sub>	Me
406	CH <sub>2</sub> CH <sub>2</sub> OCar	NO <sub>2</sub>	Et
407	NO <sub>2</sub>	H	Me
408	NO <sub>2</sub>	H	CH <sub>2</sub> OH
409	NO <sub>2</sub>	H	CH <sub>2</sub> OAc
410	NO <sub>2</sub>	H	CH <sub>2</sub> OBn
411	NO <sub>2</sub>	H	CH <sub>2</sub> OBz
412	NO <sub>2</sub>	H	CH <sub>2</sub> SH
413	NO <sub>2</sub>	H	CH <sub>2</sub> SO <sub>2</sub> Me
414	NO <sub>2</sub>	H	CH <sub>2</sub> SO <sub>2</sub> Et
415	NO <sub>2</sub>	H	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
416	NO <sub>2</sub>	H	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
417	NO <sub>2</sub>	H	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
418	NO <sub>2</sub>	H	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
419	NO <sub>2</sub>	H	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
420	NO <sub>2</sub>	H	CH <sub>2</sub> SO <sub>2</sub> Pr
421	NO <sub>2</sub>	H	CH <sub>2</sub> SO <sub>2</sub> iPr
422	NO <sub>2</sub>	H	CH <sub>2</sub> SO <sub>2</sub> Bu
423	NO <sub>2</sub>	H	CH <sub>2</sub> SO <sub>2</sub> Pn
424	NO <sub>2</sub>	H	CH <sub>2</sub> SO <sub>2</sub> Hx
425	NO <sub>2</sub>	H	CH <sub>2</sub> OCar
426	NO <sub>2</sub>	H	CH <sub>2</sub> NHCO <sub>2</sub> Me
427	NO <sub>2</sub>	H	CH <sub>2</sub> NHC(S)OMe
428	NO <sub>2</sub>	H	Et
429	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> OH
430	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> OAc
431	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> OBn
432	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> OBz
433	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SH
434	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
435	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
436	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
437	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
438	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
439	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
440	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
441	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pr

442	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
443	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Bu
444	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pn
445	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Hx
446	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> OCar
447	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> NHCO <sub>2</sub> Me
448	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> NHC(S)OMe
449	NO <sub>2</sub>	H	Pr
450	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
451	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
452	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
453	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
454	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
455	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OH)CH <sub>3</sub>
456	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OAc)CH <sub>3</sub>
457	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OBz)CH <sub>3</sub>
458	NO <sub>2</sub>	H	CH <sub>2</sub> CH(SH)CH <sub>3</sub>
459	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OH)CH <sub>2</sub> OMe
460	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> OMe
461	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> OMe
462	NO <sub>2</sub>	H	CH <sub>2</sub> CH(SH)CH <sub>2</sub> OMe
463	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OH)CH <sub>2</sub> Cl
464	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> Cl
465	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> Cl
466	NO <sub>2</sub>	H	CH <sub>2</sub> CH(SH)CH <sub>2</sub> Cl
467	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OH)CH <sub>2</sub> F
468	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
469	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
470	NO <sub>2</sub>	H	CH <sub>2</sub> CH(CH <sub>3</sub> )SO <sub>2</sub> Et
471	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
472	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
473	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
474	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
475	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
476	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pr
477	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
478	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Bu

479	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pn
480	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Hx
481	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NHCO <sub>2</sub> Me
482	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NHC (S) OMe
483	NO <sub>2</sub>	H	Bu
484	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
485	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
486	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
487	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
488	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
489	NO <sub>2</sub>	H	CH <sub>2</sub> CH (OH) CH <sub>2</sub> CH <sub>3</sub>
490	NO <sub>2</sub>	H	CH <sub>2</sub> CH (OAc) CH <sub>2</sub> CH <sub>3</sub>
491	NO <sub>2</sub>	H	CH <sub>2</sub> CH (OBz) CH <sub>2</sub> CH <sub>3</sub>
492	NO <sub>2</sub>	H	CH <sub>2</sub> CH (SH) CH <sub>2</sub> CH <sub>3</sub>
493	NO <sub>2</sub>	H	CH <sub>2</sub> CH (OH) CH <sub>2</sub> CH <sub>2</sub> OMe
494	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
495	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
496	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
497	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
498	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
499	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
500	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
501	NO <sub>2</sub>	H	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
502	NO <sub>2</sub>	H	Pn
503	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> OH
504	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> OAc
505	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> OBn
506	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> OBz
507	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> SH
508	NO <sub>2</sub>	H	CH <sub>2</sub> CH (OH) CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
509	NO <sub>2</sub>	H	CH <sub>2</sub> CH (OAc) CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
510	NO <sub>2</sub>	H	CH <sub>2</sub> CH (OBz) CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
511	NO <sub>2</sub>	H	CH <sub>2</sub> CH (SH) CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
512	NO <sub>2</sub>	H	CH <sub>2</sub> CH (OH) CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OMe
513	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> Me
514	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> Et
515	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH

516	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
517	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
518	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
519	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
520	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> iPr
521	NO <sub>2</sub>	H	Hx
522	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>6</sub> OH
523	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>6</sub> OAc
524	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>6</sub> OBn
525	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>6</sub> OBz
526	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>6</sub> SH
527	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
528	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
529	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
530	NO <sub>2</sub>	H	CH <sub>2</sub> CH(SH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
531	NO <sub>2</sub>	H	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OMe
532	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> Me
533	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> Et
534	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
535	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
536	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
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539	NO <sub>2</sub>	H	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> iPr
540	NO <sub>2</sub>	Me	Me
541	NO <sub>2</sub>	Me	CH <sub>2</sub> OH
542	NO <sub>2</sub>	Me	CH <sub>2</sub> OAc
543	NO <sub>2</sub>	Me	CH <sub>2</sub> OBn
544	NO <sub>2</sub>	Me	CH <sub>2</sub> OBz
545	NO <sub>2</sub>	Me	CH <sub>2</sub> SH
546	NO <sub>2</sub>	Me	CH <sub>2</sub> SO <sub>2</sub> Me
547	NO <sub>2</sub>	Me	CH <sub>2</sub> SO <sub>2</sub> Et
548	NO <sub>2</sub>	Me	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
549	NO <sub>2</sub>	Me	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
550	NO <sub>2</sub>	Me	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
551	NO <sub>2</sub>	Me	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
552	NO <sub>2</sub>	Me	CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH

553	NO <sub>2</sub>	Me	CH <sub>2</sub> SO <sub>2</sub> Pr
554	NO <sub>2</sub>	Me	CH <sub>2</sub> SO <sub>2</sub> iPr
555	NO <sub>2</sub>	Me	CH <sub>2</sub> SO <sub>2</sub> Bu
556	NO <sub>2</sub>	Me	CH <sub>2</sub> SO <sub>2</sub> Pn
557	NO <sub>2</sub>	Me	CH <sub>2</sub> SO <sub>2</sub> Hx
558	NO <sub>2</sub>	Me	CH <sub>2</sub> OCar
559	NO <sub>2</sub>	Me	CH <sub>2</sub> NHCO <sub>2</sub> Me
560	NO <sub>2</sub>	Me	CH <sub>2</sub> NHC (S) OMe
561	NO <sub>2</sub>	Me	Et
562	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> OH
563	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> OAc
564	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> OBn
565	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> OBz
566	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SH
567	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
568	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
569	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
570	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
571	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
572	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
573	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
574	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pr
575	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
576	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Bu
577	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pn
578	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Hx
579	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> OCar
580	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> NHCO <sub>2</sub> Me
581	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> NHC (S) OMe
582	NO <sub>2</sub>	Me	Pr
583	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
584	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
585	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
586	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
587	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
588	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OH) CH <sub>3</sub>
589	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OAc) CH <sub>3</sub>

590	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OBz) CH <sub>3</sub>
591	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (SH) CH <sub>3</sub>
592	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OH) CH <sub>2</sub> OMe
593	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OAc) CH <sub>2</sub> OMe
594	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OBz) CH <sub>2</sub> OMe
595	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (SH) CH <sub>2</sub> OMe
596	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OH) CH <sub>2</sub> Cl
597	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OAc) CH <sub>2</sub> Cl
598	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OBz) CH <sub>2</sub> Cl
599	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (SH) CH <sub>2</sub> Cl
600	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OH) CH <sub>2</sub> F
601	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
602	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
603	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (CH <sub>3</sub> ) SO <sub>2</sub> Et
604	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
605	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
606	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
607	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
608	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
609	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pr
610	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
611	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Bu
612	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Pn
613	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Hx
614	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NHCO <sub>2</sub> Me
615	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NHC (S) OMe
616	NO <sub>2</sub>	Me	Bu
617	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
618	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
619	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
620	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
621	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
622	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OH) CH <sub>2</sub> CH <sub>3</sub>
623	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OAc) CH <sub>2</sub> CH <sub>3</sub>
624	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OBz) CH <sub>2</sub> CH <sub>3</sub>
625	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (SH) CH <sub>2</sub> CH <sub>3</sub>
626	NO <sub>2</sub>	Me	CH <sub>2</sub> CH (OH) CH <sub>2</sub> CH <sub>2</sub> OMe

627	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Me
628	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> Et
629	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
630	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
631	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
632	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
633	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
634	NO <sub>2</sub>	Me	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SO <sub>2</sub> iPr
635	NO <sub>2</sub>	Me	Pn
636	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> OH
637	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> OAc
638	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> OBn
639	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> OBz
640	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> SH
641	NO <sub>2</sub>	Me	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
642	NO <sub>2</sub>	Me	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
643	NO <sub>2</sub>	Me	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
644	NO <sub>2</sub>	Me	CH <sub>2</sub> CH(SH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
645	NO <sub>2</sub>	Me	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OMe
646	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> Me
647	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> Et
648	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
649	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
650	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
651	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
652	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
653	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>5</sub> SO <sub>2</sub> iPr
654	NO <sub>2</sub>	Me	Hx
655	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> OH
656	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> OAc
657	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> OBn
658	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> OBz
659	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> SH
660	NO <sub>2</sub>	Me	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
661	NO <sub>2</sub>	Me	CH <sub>2</sub> CH(OAc)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
662	NO <sub>2</sub>	Me	CH <sub>2</sub> CH(OBz)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
663	NO <sub>2</sub>	Me	CH <sub>2</sub> CH(SH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>

664	NO <sub>2</sub>	Me	CH <sub>2</sub> CH(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OMe
665	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> Me
666	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> Et
667	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH
668	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OAc
669	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBz
670	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OBn
671	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
672	NO <sub>2</sub>	Me	(CH <sub>2</sub> ) <sub>6</sub> SO <sub>2</sub> iPr
673	iPr	NO <sub>2</sub>	Me

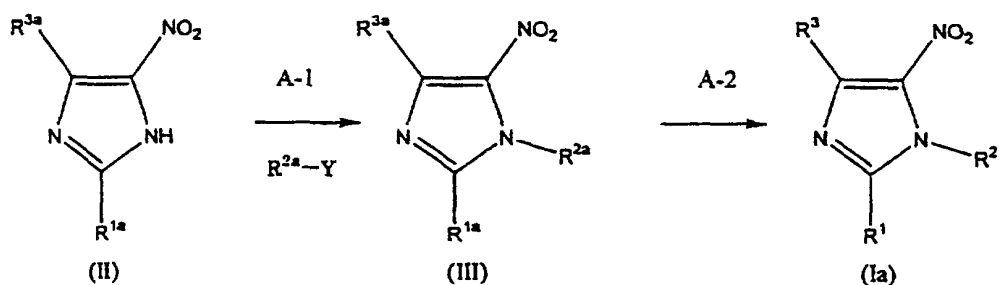
- In Table 1, Compound No. 1, 21, 23, 28, 29, 43, 50, 54, 55, 56, 57, 58, 135, 136, 137, 138, 139, 140, 141, 142, 153, 154, 155, 156, 157, 158, 159, 160, 161, 162, 163, 164, 165, 166, 167, 168, 169, 170, 175, 176, 178, 179, 180, 181, 182, 183, 184, 185, 186, 187, 191, 192, 193, 194, 195, 196, 197, 198, 209, 210, 268, 290, 291, 292, 293, 294, 295, 296, 308, 309, 316, 324, 401, 402, 403, 404, 405, 406, 459, 460, 461, 462, 493, 562, 568 or 673 is preferred; Compound No. 28, 29, 43, 54, 135, 136, 137, 138, 139, 140, 141, 142, 157, 158, 159, 160, 161, 162, 163, 175, 177, 178, 179, 180, 181, 182, 183, 191, 196, 197, 198, 290, 291, 292, 293, 294, 295, 296, 402, 459 or 673 is more preferred;
- 43) 4-(2-nitro-1H-imidazol-1-yl)ethylmorpholine (general name: Nimorazole),
- 135) 1,2-dimethyl-5-nitro-1H-imidazole (general name: Dimetridazole),
- 157) 2-(2-methyl-5-nitroimidazol-1-yl)ethanol (general name: Metronidazole),
- 163) 1-(2-ethylsulfonyl-ethyl)-2-methyl-5-nitroimidazole (general name: Tinidazole),
- 176) methyl (2-(2-methyl-5-nitroimidazol-1-yl)ethylthio-carbamate (general name: Carnidazole),
- 183) 1-(2-methyl-5-nitroimidazol-1-yl)propan-2-ol (general name: Secnidazole),
- 191) 1-chloro-3-methyl-5-nitroimidazol-1-yl)propan-2-ol

(general name: Ornidazole),  
402) 2-carbamoyloxymethyl-1-methyl-5-nitro-1H-imidazole  
(general name: Ronidazole),  
459)  $\alpha$ -methoxymethyl-2-nitroimidazole-1-ethanol (general  
5 name: Misonidazole) or  
673) 1-methyl-2-(1-methylethyl)-5-nitroimidazole (general  
name: Ipronidazole) is further more preferred; and  
157) 2-(2-methyl-5-nitroimidazol-1-yl)ethanol (general name:  
Metronidazole) or  
10 163) 1-(2-ethylsulfonylethyl)-2-methyl-5-nitroimidazole  
(general name: Tinidazole) is most preferred.

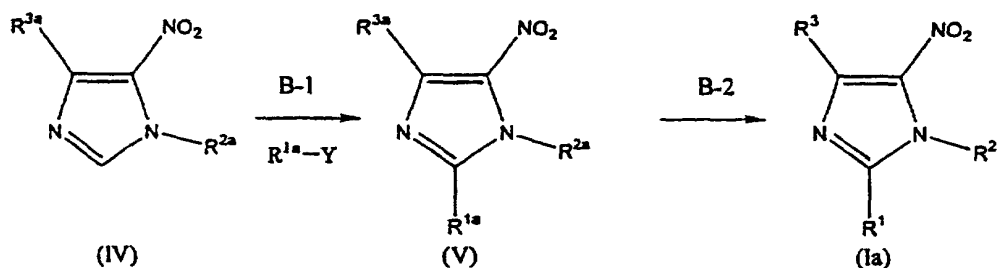
Detailed description of a preferred embodiment of the  
invention

15 The nitroimidazole derivative contained in the external  
preparation of the present invention is a known compound or  
can be also obtained by the following Process A to Process C.

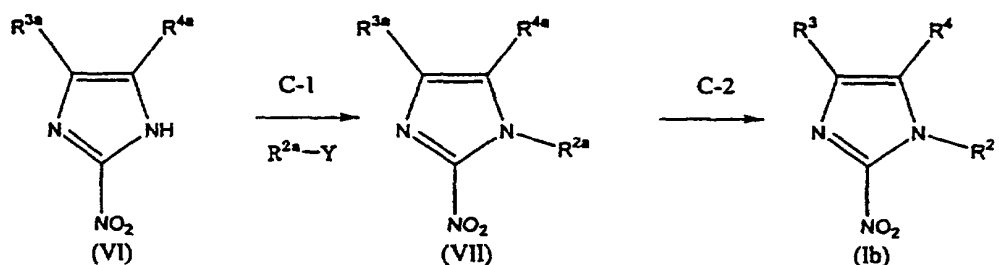
### Process A



### Process B



### Process C



In the above Processes A to Process C, R<sup>1</sup> to R<sup>4</sup> represent the same meanings as defined above and R<sup>3</sup> and R<sup>4</sup> are the same or different and represent a hydrogen atom or an optional organic group. R<sup>1a</sup> represents the above R<sup>1</sup> or a group in which a functional group on R<sup>1</sup> is appropriately protected, if necessary, according to a well-known method (for example, "Protective Groups in Organic Synthesis", Greene, T.W.; Wuts, P.G.M. John Wiley & Sons; New York, 1999, etc.) and the functional group on R<sup>1</sup> is substituted, if necessary, to an appropriate functional group capable of obtaining a desired functional group by a well-known substitution reaction. R<sup>2a</sup> represents the above R<sup>2</sup> or a group in which a functional group on R<sup>2</sup> is appropriately protected,

if necessary, according to a well-known method and the functional group on  $R^2$  is substituted, if necessary, to an appropriate functional group capable of obtaining a desired functional group by a well-known substitution reaction.  $R^{3a}$  represents the above  $R^3$  or a group in which a functional group on  $R^3$  is appropriately protected, if necessary, according to a well-known method and the functional group on  $R^3$  is substituted, if necessary, to an appropriate functional group capable of obtaining a desired functional group by a well-known substitution reaction.  $R^{4a}$  represents the above  $R^4$  or a group in which a functional group on  $R^4$  is appropriately protected, if necessary, according to a well-known method and the functional group on  $R^4$  is substituted, if necessary, to an appropriate functional group capable of obtaining a desired functional group by a well-known substitution reaction. Y represents a group to be eliminated.

In the following, respective steps of Process A to Process C are described in more detail.

(Process A)  
(Step A-1)

The present step is to prepare Compound (III) by reacting Compound (II), which is well-known or can be easily obtained from the well-known compound with  $R^{2a}$ -Y in the presence or absence of a base catalyst in an inert solvent.

The solvent employable here may include, for example, water; aliphatic hydrocarbons such as hexane, heptane, ligroin, petroleum ether, etc.; aromatic hydrocarbons, such as benzene, toluene, xylene, etc.; halogenated hydrocarbons such as methylene chloride, chloroform, carbon tetrachloride, dichloroethane, chlorobenzene, dichlorobenzene, etc.; esters such as ethyl formate, ethyl acetate, propyl acetate, butyl acetate, diethyl carbonate, etc.; ethers, such as diethyl ether, diisopropyl ether, tetrahydrofuran, dioxane, dimethoxyethane, diethylene glycol dimethyl ether, etc.; ketones, such as acetone, methyl ethyl ketone, methyl isobutyl ketone, isophorone, cyclohexanone, etc.; nitro

compounds, such as nitroethane, nitrobenzene, etc.; nitriles, such as acetonitrile, isobutyronitrile, etc.; amides, such as formamide, N,N-dimethylformamide, N,N-dimethylacetamide, N-methyl-2-pyrrolidone, N-methylpyrrolidinone,  
5 hexamethylphosphoric triamide, etc; sulfoxides, such as sulfolan, etc.; and pyridines, preferably water and the pyridine.

The base catalyst employable here may include, for example, alkali metal hydroxides, such as sodium hydroxide,  
10 potassium hydroxide, etc.; alkali metal carbonates, such as sodium carbonate, potassium carbonate, etc; alkali metal alkoxides, such as sodium methoxide, sodium ethoxide, etc.; organic bases, such as triethylamine, pyridine, dimethylaminopyridine, etc.; and ammonia water, preferably  
15 the alkali metal hydroxides.

A reaction temperature varies depending on the raw material compound, the solvent, the reagent and the base catalyst, and is usually 0°C to 200°C, preferably 20°C to 130°C.

20 A reaction time varies depending on the raw material compound, the solvent, the reagent, the base catalyst and the reaction temperature, and is usually 10 minutes to 3 days, preferably 1 to 10 hours.

After completion of the reaction, the desired Compound  
25 (III) of the present reaction is obtained, for example, by neutralizing the reaction mixture, concentrating the reaction mixture, adding an organic solvent immiscible with water such as ethyl acetate, washing with water, separating an organic layer or an aqueous layer containing the desired compound and  
30 distilling off the solvent.

The compound, thus obtained, can be further purified, if necessary, by a conventional method, for example, recrystallization and silica gel column chromatography.

In the case where the compound, thus obtained, is the  
35 desired Compound (Ia) and deprotection and conversion of a functional group are not required, Compound (Ia) can be obtained without conducting Step A-2 described below.

(Step A-2)

5 The present step is to prepare Compound (Ia) by carrying out a substitution reaction of Compound (III) which is well-known or is obtained in (A-1), if necessary, in an inert solvent, subsequently or concurrently carrying out the deprotection reaction if necessary.

10 The substitution reaction of the present step varies depending on the desired substituents and is not particularly limited so long as it is a reaction in which the desired functional group is obtained and is carried out according to a method described in references (for example, "Aliphatic Nucleophilic Substitution", Hartshorn, Cambridge University Press: Cambridge (1973), Chem. Soc. Rev., 19, 83 (1990), Carbocation Chem., 1, 121 (1989), etc.).

15 The deprotection reaction of the present step varies depending on the protective group and is not particularly limited so long as it is the reaction in which the desired functional group is obtained and is carried out according to a method described in references (for example, "Protective Groups in Organic Synthesis", Greene, T.W.; Wuts, P.G.M. John Wiley & Sons; New York, 1999, etc.).

(Process B)

25 (Step B-1)

The present step is to prepare Compound (V) by reacting Compound (IV) which is publicly known or can be easily obtained from the publicly known compound with  $R^{1a}-Y$  in the presence or absence of a base catalyst in an inert solvent.

30 The solvent employable here may include, for example, water; aliphatic hydrocarbons such as hexane, heptane, ligroin and petroleum ether; aromatic hydrocarbons such as benzene, toluene and xylene; halogenated hydrocarbons such as methylene chloride, chloroform, carbon tetrachloride, dichloroethane, chlorobenzene and dichlorobenzene; esters such as ethyl formate, ethyl acetate, propyl acetate, butyl acetate and diethyl carbonate; ethers such as diethyl ether,

diisopropyl ether, tetrahydrofuran, dioxane, dimethoxyethane and diethylene glycol dimethyl ether; ketones such as acetone, methyl ethyl ketone, methyl isobutyl ketone, isophorone and cyclohexanone; nitro compounds such as nitroethane and  
5 nitrobenzene; nitriles such as acetonitrile and isobutyronitrile; amides such as formamide, N,N-dimethylformamide, N,N-dimethylacetamide, N-methyl-2-pyrrolidone, N-methylpyrrolidinone and hexamethylphosphoric triamide; sulfoxides such as sulfolan; and pyridines,  
10 preferably water and the pyridine.

The base catalyst employable here may include, for example, alkali metal hydroxides such as sodium hydroxide and potassium hydroxide; alkali metal carbonates such as sodium carbonate and potassium carbonate; alkali metal alkoxides  
15 such as sodium methoxide and sodium ethoxide; organic bases such as triethylamine, pyridine and dimethylaminopyridine; and ammonia water, preferably the alkali metal hydroxides.

The reaction temperature varies depending on the material compound, the solvent, the reagent and the base  
20 catalyst and is usually 0°C to 200°C, preferably 20°C to 130°C.

The reaction time varies depending on the material compound, the solvent, the reagent, the base catalyst and the reaction temperature and is usually 10 minutes to 3 days, preferably 1 to 10 hours.

25 After completion of the reaction, the desired Compound (V) of the present reaction is obtained, for example, by neutralizing the reaction mixture, concentrating the reaction mixture, adding an organic solvent immiscible with water such as ethyl acetate, washing with water, separating an organic  
30 layer or an aqueous layer containing the desired compound and distilling off the solvent.

The compound thus obtained can be further purified, if necessary, by a conventional method, for example, recrystallization and silica gel column chromatography.

35 In the case where the compound thus obtained is the desired Compound (Ia) and deprotection and conversion of a functional group are not required, Compound (Ia) can be

obtained without conducting Step B-2 described below.

(Step B-2)

5 The present step is to prepare Compound (Ia) by carrying out the substitution reaction of Compound (V) which is publicly known or is obtained in (B-1), if necessary, in an inert solvent, subsequently or concurrently carrying out the deprotection reaction if necessary.

10 The present step is carried out in the same manner as in Step A-2.

(Process C)

(Step C-1)

15 The present step is to prepare Compound (VII) by reacting Compound (VI) which is publicly known or can be easily obtained from the publicly known compound with  $R^{2a}-Y$  in the presence or absence of a base catalyst in an inert solvent.

20 The present step is carried out in the same manner as in Step A-1.

The compound, thus obtained, can be further purified, if necessary, by a conventional method, for example, recrystallization and silica gel column chromatography.

25 In the case where the compound thus obtained is the desired Compound (Ib) and deprotection and conversion of a functional group are not required, Compound (Ib) can be obtained without conducting Step C-2 described below.

(Step C-2)

30 The present step is to prepare Compound (Ib) by carrying out the substitution reaction of Compound (VII) which is publicly known or is obtained in (C-1), if necessary, in an inert solvent, subsequently or concurrently carrying out deprotection reaction if necessary.

35 The present step is carried out in the same manner as in Step A-2.

Moreover, the compound of the present invention can be

obtained in accordance with known methods for example, a production method of metronidazole is disclosed by Jacob, et al. (US Patent No. 2,944,061) and a production method of tinidazole is disclosed by Butler, et al. (US Patent No. 3,376,311); and, for example, a production method of a derivative having a dihydropyridine ring is disclosed by Gorlitzer, et al. (Pharmazie (1999), 54(12), 889-892), a production method of a carbamate derivative is disclosed by Hay, et al. (Bioorg. Med. Chem. Lett. (1999), 9(15), 2237-2242), a production method of a derivative having an isoquinoline ring is disclosed by Parveen, et al. (Bioorg. Med. Chem. Lett. (1999), 9(15), 2031-2036), a production method of a derivative having a pyrrole ring is disclosed by Anadlu, et al. (Eur. J. Med. Chem. (1999), 34(3), 275-278), a production method of a derivative having a benzene ring substituted with a carboxyl group is disclosed by Everett, et al. (Bioorg. Med. Chem. Lett. (1999), 9(9), 1267-1272), a production method of a derivative having a halogen-substituted benzene ring and a derivative having a pyridine ring is disclosed by Shafiee, et al. (J. Heterocycl. Chem. (1998), 35(3), 607-610), a production method of a derivative having an arylcarbonyloxy group is disclosed by Bowden, et al. (Eur. J. Med. Chem. (1997), 32(12), 995-1000), a production method of a derivative having a dioxolane ring is disclosed by Baji, et al. (Eur. J. Chem. (1997), 32(7-8), 637-650), a production method of a derivative having a hydroxyaryl group is disclosed by Arrendondo, et al. (Bioorg. Med. Chem. Lett. (1996), 6(15), 1781-1784), a production method of a derivative having a phenylamide group is disclosed by Shafiee, et al. (J. Sci. Islamic Repub. Iran (1995), 6(1), 25-8), a production method of a derivative having an alkylthio group is disclosed by Rao, et al. (J. Chem. Soc. Perkin Trans. 1 (1994), (17), 2399-2402), a production method of a derivative having a hydroxyalkylaryl group is disclosed by Furlan, et al. (European Unexamined Patent Publication No. EP535528), a production method of a derivative having a  $\beta$ -lactam ring is disclosed by Bertola, et al. (European Unexamined Patent

Publication No. EP490450), a production method of an aminoalkylphenylacyl derivative is disclosed by Bundgaard, et al. (International Unexamined Patent Publication No. WO90/08128), a production method of a derivative having an aralkylcarbonyloxy group is disclosed by Rao, et al. (Indian J. Chem., Sect. B (1990), 29B(11), 1034-1040), a production method of an N-substituted aminomethylbenzoate derivative is disclosed by Jansen, et al. (Int. J. Pharm. (1990), 58(2), 143-153), a production method of a derivative having a pyridine ring is disclosed by Alcalde, et al. (Farmaco (1989), 44(11), 1095-1097), a production method of metronidazole and secnidazole accompanying a nitro group transfer reaction is disclosed by Bufo, et al. (US Patent No. 4,925,591), a production method of dimetridazole using dimethyl sulfate is disclosed by Bufo, et al. (US Patent No. 4,925,952), a production method of metronidazole.secnidazole using alkylene sulfate is disclosed by Bonnamas, et al. (US Patent No. 4,925,949), a production method of a derivative having an alkylcarbonyloxy group is disclosed by Johansen, et al. (Int. J. Pharm. (1986), 32(2-3), 199-206), a production method of a derivative having a dextran ring is disclosed by Vermeersch, et al. (Bull. Soc. Chim. Belg. (1985), 94(8), 591-596), a production process of an L-cysteine derivative is disclosed by Reiner (European Unexamined Patent Publication No. 140395), a production method of an ester derivative of an amino acid residue is disclosed by Cho (European Unexamined Patent Publication No. 127274), a production method of a derivative having a hydroxyalkylaryl group is disclosed by Tessitore (European Unexamined Patent Publication No. 11657), a production method of a derivative having a hydroxyalkylaryl group is disclosed by Scalesiani (European Unexamined Patent Publication No. EP103100), a production method of an ester derivative of an amino acid residue is disclosed by Bundgaard, et al. (Int. J. Pharm. (1984), 18(1-2), 67-77), a production method of an ester derivative of a dimethylglycine residue is disclosed by Thorbek, et al. (European Unexamined Patent Publication No. 96870), a production method of a 2-

nitroimidazole derivative having an aryloxy ring is disclosed by Hofheinz (European Unexamined Patent Publication No.

EP81719), a production method of a platinum salt derivative is disclosed by Bales, et al. (J. Chem. Soc., Chem. Comm.

5 (1983), (8), 432-433), a production method of a retinoic acid derivative is disclosed by Whitefield, et al. (UK Unexamined Patent Publication No. 2097783), a production method of a

derivative having a hydroxyalkylaryl group is disclosed by

Bononi (US Patent No. 4,463,012), a production method of a

10 derivative having an alkylsulfonylphenyloxy group and a production method of a derivative having alkylthiophenyloxy group are disclosed by Winkelmann, et al. (UK Unexamined

Patent Publication No.s 1541280 and 1590974), a production method of a derivative having an acetamide-substituted

15 phenyloxy group is disclosed by Winkelmann, et al. (Arzneim.-Forsch. (1978), 28(5), 739-49), a production method of a derivative having a phenylcarbamate group and a production

method of a derivative having a hydrazinecarboxyl group are disclosed by Cavalleri, et al. (J. Med. Chem. (1978), 21(8),

20 781-4), a production method of a derivative having a hydrazine group is disclosed by Winkelmann, et al. (Arzneim.-Forsch. (1977), 27(12), 2251-63), a production method of a

2-nitroimidazole derivative having a phenylcarbamate group is

disclosed by Cavalleri, et al. (US Patent No. 4,113,953), a

25 production method of a 2-nitroimidazole derivative having an aluren group is disclosed by Cavalleri, et al. (J. Med. Chem. (1977), 20(11), 1522-5), a production method of a derivative

having a substituted phenoxy group is disclosed by Winkelmann, et al. (US Patent No. 4,031,232), a production method of a 2-

30 nitroimidazole derivative having a hydroxy group is disclosed by Assandri, et al. (UK Unexamined Patent Publication No. 1480192), a production method of a derivative having an

alkylcarbonyloxy group is disclosed by Hoffer (UK Unexamined Patent Publication No. 1453417), a production method of a

35 derivative having a quinoline ring is disclosed by Kreider, et al. (US Patent No.s 3,910,925 and 3,828,056), a production method of dimetridazole using formic acid as a solvent is

disclosed by Frank, et al. (UK Unexamined Patent Publication No. 1493496), a production method of metronidazole using alkylene oxide is disclosed by Frank, et al. (UK Unexamined Patent Publication No. 1481349), a production method of a derivative having a phenylamide group is disclosed by Heeres, et al. (US Patent No. 3,928,374), a production method of ornidazole using epichlorohydrin is disclosed by Hoffer, et al. (J. Med. Chem. (1974), 17(9), 1019-20), a production method of 1-methyl-5-isopropyl-2-nitroimidazole is disclosed by Martin, et al. (US Patent No. 3,828,064), a production method of a derivative having a carboxyimidamide group is disclosed by Rufer, et al. (US Patent No. 3,966,732), a production methods of a derivative having a carbamate group and a 2-nitromidazole derivative are disclosed by Challier, et al. (UK Unexamined Patent Publication No. 1352288), a production method of metronidazole using ethylene oxide and sulfuric acid is disclosed by Muhlbrod (UK Unexamined Patent Publication No. 1301225), a production method of a 2-nitromidazole derivative having a phenylhydrazone group or alkenyl group is disclosed by Cavalleri, et al. (J. Heterocycl. Chem. (1972), 9(5), 979-84), a production method of a derivative having a carboxyimidyl group is disclosed by Papaioannou (US Patent No. 3,694,452), a production method of a derivative having an alkenyl group is disclosed by Hoffer, et al. (US Patent No. 3,652,579), a production method of a 2-benzyl derivative is disclosed by Hoff, et al. (UK Unexamined Patent Publication No. 1336228), a production method of secnidazole by nitration using nitric acid and its following hydrolysis is disclosed by Jeanmart, et al. (UK Unexamined Patent Publication No. 1278757), a production method of secnidazole and 1-(2-methyl-5-nitroimidazol-1-yl)2-propanone is disclosed by Jeanmart, et al. (UK Unexamined Patent Publication No. 1278758), a production method of 1,5-dimethyl-2-nitroimidazole and 1,4,5-trimethyl-2-nitroimidazole is disclosed by Lancini, et al. (UK Unexamined Patent Publication No. 1114154), a production method of a derivative having a nitroso group is disclosed by Kulsa, et

al. (US Patent No. 3,711,495), a production method of a derivative having a hydroxy group is disclosed by Chemerda, et al. (US Patent No. 3,584,007), a production method of a 2-nitroimidazole derivative substituted with an alkyl group is disclosed by Lancini, et al. (South African Unexamined Patent Publication No. 6905670), a production method of a derivative having a sulfonyl group (including tinidazole) is disclosed by Miller, et al. (J. Med. Chem. (1970), 13(5), 849-52), a production method of a 2-nitroimidazole derivative having a hydroxy group is disclosed by Lancini, et al. (UK Unexamined Patent Publication No. 1229170), a production method of a derivative having an alkyl halide group is disclosed by Kajfez, et al. (Farm. Glas. (1969), 25(2), 49-54), and a production method of 4-iodo-1,2-dimethyl-5-nitroimidazole is disclosed by Hoffer, et al. (J. Heterocycle Chem. (1966), 3(4), 454-8)).

Examples of administration forms of the external preparation for skin diseases of the present invention include ointment, cream, lotion, moisturized patch or moisture-free patch, shampoo, gel, rinse, face lotion, milky lotion, paste, shaving cream, foundation, cologne, pack, semi-solid, solid or liquid. These preparations can be produced in accordance with routine methods, for example, as described below using, as necessary, additives such as antioxidants (including carboxylic acids such as ascorbic acid and citric acid; and phenols such as tocopherol and dibutylhydroxytoluene), antiseptics (including carboxylic acids such as dehydroacetic acid, salicylic acid and disodium edetate; and phenols such as ethyl paraoxybenzoate, methyl paraoxybenzoate, isopropyl paraoxybenzoate and thymol), wetting agents (including glycols such as glycerin, propylene glycol, dipropylene glycol and 1,3-butylene glycol; organic salts such as hyaluronic acid; and amides such as urea), consistency agents (including polymer compounds such as polyethylene glycol; and celluloses such as carboxymethyl cellulose sodium and carboxypropyl cellulose), buffers (including organic acids such as citric acid, lactic acid and

tartaric acid; inorganic acids such as hydrochloric acid and boric acid; salts such as sodium dihydrogen phosphate and sodium citrate; organic bases such as triethanolamine; and inorganic bases such as sodium hydroxide and potassium hydroxide), adsorbents (including water-containing aluminum silicates such as kaolin and bentonite; and inorganic bases such as magnesium hydroxide-aluminum hydroxide co-precipitate and aluminum hydroxide), bases (including organic substances such as white petrolatum, Tween 60, Tween 80, liquid paraffin, beeswax, petrolatum, castor oil, silicone oil, hydrogenated castor oil, natural rubber, coconut oil fatty acid diethanolamide, polyoxyethylene hydrogenated castor oil, natural rubber latex and 1,3-pentadiene copolymer resin; polymer compounds such as polybutene, synthetic rubber SBR, polyethylene glycol monostearate, polyoxyethylene glycol monostearate, polyoxyethylene cetostearyl ether, polyoxyethylene oleylcetyl ether, silicon, starch grafted acrylate 300, sodium polyacrylate, methacrylic acid-n-butyl acrylate copolymer and carboxyvinyl polymer; fatty acids such as stearic acid; alcohols such as cetanol and myristyl alcohol, and fatty acid esters such as octadodecyl myristate, isopropyl myristate and cetyl octanoate), solvents (including carbohydrates such as ethanol, isopropanol, 1,3-butylene glycol, n-octadecylalcohol, crotamiton and caprylic/capric acid triglyceride), stabilizers (including inorganic salts such as sodium metaphosphate, zinc oxide and titanium oxide; and organic salts such as sodium polyoxyethylene lauryl sulfate ether sulfate and sodium lauryl sulfate), adhesives (including polymer compounds such as sodium polyacrylate and dipropylene glycol), emulsifiers (including carbohydrates such as sorbitan monooleate, polyoxyethylene sorbitan monooleate, D-sorbitol, polyglycerin monolaurate and sodium polyoxyethylene lauryl ether sulfate), surfactants (including polymer compounds such as polyglycerin monolaurate and polyoxyethylene oleyl alcohol ether), squalane, diluent, Span 60, Span 80, gelatin, propylparaben, methylparaben, lauryldimethyl-aminoacetate betaine, coconut oil fatty acid

diethanol amide, N-[alkyl(12,14)oxy-2-hydroxypropyl]-L-arginine hydrochloride, silicone oil, jojoba oil, and fragrance.

Any fragrances can be used provided they can be generally used in foods, cosmetics, pharmaceuticals and so forth. Examples of naturally-derived fragrances include those obtained from plants such as rose, lavender and orange, and those obtained from animals such as musk oil (musk) obtained from musk deer and castorium (castor oil) obtained from beavers. Examples of synthetic fragrances include limonene,  $\beta$ -caryophyllene, farnesol, citral,  $\gamma$ -undecalactone, indole and rilal.

Ointments are produced by, for example, heating and stirring an active ingredient and base, heating and dispersing, followed by cooling to the room temperature while stirring.

Creams are produced by, for example, first producing a base while heating and stirring, adding an active ingredient itself or a solution containing the active ingredient while heating and stirring, and cooling the resulting emulsion to the room temperature.

Lotions are produced by, for example, adding the active ingredient itself or a solution containing the active ingredient to an oily base or mixed base consisting of an oily base melted by heating and an aqueous base while stirring and heating, and then adding an aqueous base and cooling the resulting liquid to the room temperature.

Moisturized patches are produced by, for example, adding an additive to a mixed base consisting of an oily base melted by heating and an aqueous base while stirring, adding an active ingredient or a solution containing the active ingredient to the mixture while heating with stirring, rolling out the resulting paste onto a non-woven fabric and cutting to an appropriate size.

Moisture-free patches are produced by, for example, adding an active ingredient or a solution containing the active ingredient to a mixed base consisting of an oily base

melted by heating while heating and stirring, adding this to a mixture of synthetic resin that has been melted by heating while stirring, rolling out the resulting paste onto a non-woven or woven fabric and cutting to an appropriate size.

5       Gels are produced by, for example, uniformly dissolving a gel base followed by adding a hydrophilic organic solvent, adding an active ingredient, heating, dissolving and dispersing. A solvent is then added thereto while heating. Next, after neutralizing while stirring, the mixture is  
10       cooled to the room temperature.

Shampoos are produced by, for example, heating purified water, adding an active ingredient, anionic surfactant, humectant and so forth, and cationic polymer as necessary, followed by uniformly dissolving and then cooling.

15       Pastes are produced by, for example, adding fats and oils to a wax, heating to melt, adding pigment, hydrocarbon and effective ingredient and so forth, and humectant as necessary, followed by mixing uniformly and cooling.

Rinses are produced by, for example, adding aqueous  
20       ingredients such as effective ingredient, humectant and cationic surfactant to purified water followed by melting with heating. Oily components such as higher alcohols and hydrocarbons are then added thereto after melting with heating followed by stirring to obtain a uniform mixture and  
25       then cooling.

Liquids are produced by, for example, adding and mixing an effective ingredient, humectant, lower alcohol and so forth to purified water, and then adding water-soluble polymer as necessary. Liquids can also be produced by adding  
30       these to mixture of oily components such as fatty acids, fats and oils and fatty acid esters as necessary followed by melting with heating.

Soap is produced by, for example, adding alkali to heated fats and oils. Alternatively, soap is produced by  
35       adding and stirring an added lower alcohol in fats and oils followed by the addition of alkali, purified water and humectant. Polysaccharides may also be added to this and

mixed thoroughly followed by the addition of dye, fragrance and an effective ingredient followed by mixing uniformly, cooling and drying to obtain soap.

5 Milky lotions can be produced by, for example, adding an effective ingredient and humectant, etc. to purified water followed by heating to melt, adding this to oily components such as surfactant and higher alcohol that have been melted by heating, and then mixing uniformly and cooling.

10 Shaving creams can be produced by, for example, adding an active ingredient, humectant, alkali and so forth to purified water followed by heating and melting. This is then added to a mixture of necessary ingredients such as fatty acid, fatty acid esters, fats and oils and so forth that have been melted by heating followed by mixing uniformly and  
15 cooling.

Face lotions are produced by, for example, adding an active ingredient, thickener, humectant and so forth to purified water followed by the addition of a mixture of alcohol, surfactant and oily components such as fats and oils  
20 and mixing uniformly.

Foundations are produced by, for example, mixing pigments and coloring pigments of finely ground clay minerals, adding fatty acid, higher alcohols and other fats and oils and esters and mixing uniformly.

25 Colognes are produced by, for example, adding and mixing an effective ingredient, humectant, lower alcohol and so forth into purified water, adding water-soluble polymer as necessary and then adding fragrance after cooling. If necessary, colognes can also be produced by adding these to a  
30 mixture of oily ingredients such as fatty acids, fats and oils and fatty acid esters, etc. after melting by heating, and then adding fragrance after cooling.

The raw materials used in packs are completely different depending on the preparation form. If the pack is in the  
35 form of a jelly, it is produced by, for example, heating and melting an effective ingredient, humectant, alkali and so forth in purified water, adding thickener, water-soluble

polymer and so forth, followed by stirring. Next, alcohols, surfactant and so forth are added and dissolved followed by cooling.

Further, in the production of the external preparation for skin diseases of the present invention, other pharmaceutically effective ingredients may be contained therein in addition to those agents provided that they do not impair the effect of combining those agents which is dermatologically applicable. Examples of these pharmaceutically active ingredients include known refrigerants, keratolytics, cortical inhibitors, antiseborrheics, germicides, antipruritics as well as drugs that can be used for skin diseases, specific examples of which include menthol, salicylic acid, estradiol, glycyrrhizic acid, benzalkonium chloride, phenol and camphor; narcotics and antihypnotics such as ethylmorphine hydrochloride, oxycodone hydrochloride, cocaine hydrochloride, pethidine hydrochloride, methamphetamine hydrochloride, dl-methylephedrine hydrochloride, morphine hydrochloride, fentanyl citrate, levallorphan tartrate; local germicides such as povidone iodide and iodoform; enzyme preparations such as lysozyme hydrochloride, streptokinase, streptodornase trypsin and deoxyribonuclease; herbal medicines such as Lithospermi Radex extract and scopolia extract; hemorrhoid preparations such as non-viable E. coli, epidihydrocholesterin and tribenoside; and hemostyptics such as thrombin, cellulose oxide and sodium alginate.

The following provides a more detailed explanation of the present invention through its Examples and Test Examples.

[Example]

(Example 1) Ointment for external use

Prescription:

The ointment is composed of metronidazole (2 g), Tween 80 (1 g), propylene glycol (28 g) and white petrolatum (69 g).

Preparation method:

A mixture of Tween 80, propylene glycol and metronidazole was added to a white petrolatum, while heating and stirring. The mixture was dispersed with continuously stirring while heating. The dispersion was then allowed to cool slowly to about 25°C and charged in a suitable vessel.

(Example 2) Ointment for external use

Prescription:

The ointment is comprised of metronidazole (2 g), propylene glycol (5 g), polyoxyethylene glycol monostearate (4 g), liquid paraffin (10 g), white petrolatum (60 g) and distilled water (an amount making total 100 g).

Preparation method:

Distilled water, propylene glycol and metronidazole were dispersed under stirring and heating and a temperature of the dispersion was adjusted to about 70°C. Polyoxyethylene glycol monostearate, liquid paraffin and white petrolatum were melted and adjusted to about 70°C and the molten mixture was slowly added to the dispersion. The mixture was allowed to cool slowly to about 25°C while continuously stirring, and charged in a suitable vessel.

(Example 3) Cream for external use

Prescription:

The cream is comprised of metronidazole (2 g), stearic acid (5 g), polyoxyethylene cetostearyl ether (12E.O.) (0.5 g), polyoxyethylene cetostearyl ether (20E.O.) (0.5 g), cetanol (5 g), cetyl octanoate (5 g), liquid paraffin (5 g), beeswax (1 g), glycerin (5 g), 1,3-butylene glycol (5 g), triethanolamine (5 g), hydrochloric acid (2.7 g) and distilled water (an amount making total 100 g).

Preparation method:

Polyoxyethylene cetostearyl ether (20E.O.), polyoxyethylene cetostearyl ether (12E.O.), stearic acid, cetanol, cetyl octanoate, liquid paraffin and beeswax were melted at a temperature of about 70 to 75°C. Distilled water, glycerin, 1,3-butylene glycol and triethanolamine were

dissolved and kept at a temperature of about 70°C, and slowly added thereto while stirring. Subsequently, distilled water, metronidazole and hydrochloric acid were dissolved and the solution was heated to about 70°C and slowly added thereto.

- 5 The emulsion thus obtained was cooled to a temperature of about 25°C while continuously stirring. The cream thus obtained was charged in a suitable vessel.

(Example 4) Cream for external use

10 Prescription:

The cream is composed of metronidazole (1.8 g), stearic acid (2 g), glycol monostearate (12 g), polyoxyethylene glycol monostearate (3 g), polyoxyethylene cetostearyl ether (12E.O.) (1 g), polyoxyethylene cetostearyl ether (20E.O.) (1 g), cetanol (2 g), liquid paraffin (5 g), cetyl octanoate (5 g), ethyl paraoxybenzoate (0.3 g), silicone oil (1 g), beeswax (1.5 g), 1,3-butylene glycol (7 g), glycerin (5 g), sodium hydroxide (suitable amount), hydrochloric acid (suitable amount) and distilled water (an amount making total 100 g).

20 Preparation method:

To a dissolved solution of distilled water, 1,3-butylene glycol and glycerin, metronidazole was added and the hydrochloric acid was added until metronidazole was dissolved completely. The solution was heated to about 70°C and pH was made to 6.9 by the addition of sodium hydroxide. As oil phase, stearic acid, glycol monostearate, polyoxyethylene glycol monostearate, polyoxyethylene cetostearyl ether (12E.O.), polyoxyethylene cetostearyl ether (20E.O.), cetanol, liquid paraffin, cetyl octanoate, silicone, ethyl paraoxybenzoate and beeswax were melted and adjusted to temperature of about 70 to 75°C. To this oil phase mixture, the foregoing solution was slowly added while stirring. The emulsion thus obtained was cooled to a temperature of about 25°C while continuously stirring. The cream thus obtained was charged in a suitable vessel.

(Example 5) Cream for external use

Prescription:

The cream is composed of metronidazole (1.8 g), n-octadecyl alcohol (5 g), stearic acid (5 g), triethanolamine (5 g), liquid paraffin (10 g), disodium edetate (0.25 g), glycerin (10 g), thymol (0.25 g), hydrochloric acid (suitable amount) and distilled water (an amount making total 100 g).

Preparation method:

A mixture of n-octadecyl alcohol, stearic acid and liquid paraffin was melted by heating while stirring and kept at about 70°C, and metronidazole was then added thereto. While keeping a temperature at about 70°C and stirring, a dissolved mixture of distilled water, glycerin and triethanolamine was slowly added thereto. Subsequently, disodium edetate and thymol were added thereto. The emulsion thus obtained was adjusted to pH 6.8 by adding hydrochloric acid, followed by cooling to a temperature of about 25°C while continuously stirring and charged in a suitable vessel.

(Example 6) Lotion for external use

Prescription:

The lotion is composed of metronidazole (2 g), stearic acid (4 g), cetanol (1 g), polyoxyethylene cetostearyl ether (20E.O.) (1 g), triethanolamine (0.2 g), glycerin (5 g), isopropanol (10 g), and distilled water (an amount making total 100 g).

Preparation method:

Cetanol, polyoxyethylene cetostearyl ether (20E.O.), stearic acid and metronidazole were melted by heating while stirring. A molten mixture of triethanolamine, distilled water and glycerin was added thereto. The mixture was then cooled to a temperature of 40°C and isopropanol was added thereto. The mixture was cooled rapidly to a temperature of about 25°C while continuously stirring. After cooling, the mixture was charged in a suitable vessel.

(Example 7) Lotion for external use

Prescription:

The lotion is composed of metronidazole (1.8 g), n-octadecyl alcohol (1 g), cetanol (1 g), polyoxyethylene cetostearyl ether (12E.O.) (1 g), 1,3-butylene glycol (10 g), Tween 80 (1 g), sodium carboxymethyl cellulose (1 g), isopropanol (10 g), distilled water (an amount making total 100 g).

Preparation method:

A temperature of a dissolved-mixture comprising 1,3-butylene glycol and distilled water was adjusted a temperature to about 70°C while continuously stirring. N-octadecyl alcohol, cetanol and polyoxyethylene cetostearyl ether (12E.O.) were melted by heating and adjusted to about 70°C, then slowly added to the above mixture. Then, under stirring, a material obtained by mixing metronidazole, sodium carboxymethyl cellulose and Tween 80 under heating was added to the above mixture. After the resulting mixture was cooled to a temperature of about 40°C, isopropanol was slowly added thereto. The mixture was cooled to a temperature of about 25°C while stirring and charged in a suitable vessel.

(Example 8) Moisture patch

Prescription:

The patch is composed of metronidazole (2 g), kaolin (5 g), liquid paraffin (10 g), glycerin (15 g), sodium carboxymethyl cellulose (5 g), crotamiton (1.5 g), zinc oxide (2 g), Tween 80 (1 g), gelatin (5 g), sodium polyacrylate (5 g), distilled water (an amount making total 100 g).

Preparation method:

Distilled water, sodium carboxymethyl cellulose and gelatin were melted by heating and added to a mixture in which zinc oxide, sodium polyacrylate and liquid paraffin were dispersed by stirring. Kaolin was added thereto while stirring. Subsequently, metronidazole, crotamiton, glycerin and Tween 80 were mixed under stirring and heating, adjusted a temperature to about 60°C and added to the above mixture under stirring and heating. Ointment thus obtained was

applied with 1000 g/m<sup>2</sup> to an unwoven fabric and the fabric was cut into a size of 10 cm x 14 cm (280 mg of metronidazole was contained per 14 g of ointment).

5 (Example 9) Moisture patch

Prescription:

The patch is composed of metronidazole (2 g), sorbitan monooleate (0.5 g), polyoxyethylene sorbitan monooleate (0.5 g), castor oil (1 g), crotamiton (1 g), gelatin (1 g), kaolin  
10 (12 g), sodium metaphosphate (0.15 g), 1,3-butylene glycol (5 g), starch grafted acrylate 300 (1 g), sodium polyacrylate (5 g), a methacrylic acid-n-butyl acrylate copolymer (3 g), D-sorbitol solution (70%) (50 g), tartaric acid (1.5 g),  
15 titanium oxide (1 g), magnesium hydroxide-aluminum hydroxide co-precipitate (0.25 g), dibutylhydroxytoluene (0.2 g) and distilled water (an amount making total 100 g).

Preparation method:

Suitable amounts of distilled water and a D-sorbitol solution were mixed and dissolved. While continuously  
20 stirring, titanium oxide was added, and then, suitable amounts of kaolin and a D-sorbitol solution were added thereto. To the mixture was added a sodium metaphosphate solution dissolved in distilled water, then a gelatin solution dissolved in distilled water was added, and a  
25 methacrylic acid-n-butyl acrylate copolymer was further added. To the above mixture were added a dissolved mixture of sodium polyacrylate, starch grafted acrylate 300, magnesium hydroxide-aluminum hydroxide co-precipitate, 1,3-butylene glycol and castor oil, sorbitan monooleate and  
30 polyoxyethylene sorbitan monooleate, and a heated mixture of metronidazole, crotamiton and dibutylhydroxytoluene was added thereto. A finally remaining mixture of D-sorbitol solution and tartaric acid was adjusted to a temperature of 60°C and added thereto while stirring. Ointment thus obtained was  
35 applied with 1000 g/m<sup>2</sup> to an unwoven fabric and the fabric was cut into a size of 10 cm x 14 cm (280 mg of metronidazole was contained per 14 g of ointment).

(Example 10) Dry patch (plaster)

Prescription:

The patch is composed of metronidazole (2 g), liquid  
5 paraffin (8 g), dibutylhydroxytoluene (0.2 g), crotamiton (1 g),  
polyoxyethylene glycol monostearate (2 g), polyoxyethylene  
cetostearyl ether (20E.O.) (1.8 g), methacrylic acid-n  
acrylate copolymer (5 g), myristyl alcohol (8 g), natural  
rubber (20 g), synthetic rubber SBR (37 g), and polybutene  
10 (15 g).

Preparation method:

Metronidazole, dibutylhydroxytoluene and crotamiton  
were mixed under stirring and heating. Subsequently,  
polyoxyethylene glycol monostearate, polyoxyethylene  
15 cetostearyl ether (20E.O.) and myristyl alcohol were added to  
the above mixture and the resulting mixture was mixed under  
heating. The resulting mixture was continuously added to a  
molten state mixture of a natural rubber latex, a methacrylic  
acid-n-butyl acrylate copolymer and a synthetic rubber SBR  
20 while stirring. Liquid paraffin and polybutene were  
continuously added thereto while stirring. Ointment thus  
obtained was applied with 100 g/m<sup>2</sup> to an unwoven fabric or  
woven fabric. After drying, the fabric was cut into a size  
of 10 cm x 14 cm (28 mg of metronidazole was contained per  
25 1.4 g of ointment).

(Example 11) Ointment for external use

Prescription:

The ointment is composed of tinidazole (2 g), propylene  
30 glycol (28 g), cetyl octanoate (5 g), and white petrolatum  
(65 g).

Preparation method:

Propylene glycol was added to white petrolatum under  
heating and stirring. A mixed material of tinidazole and  
35 cetyl octanoate was added to the above mixture and the  
resulting mixture was heated while continuously stirring to  
disperse the material therein. Then, the dispersion was

allowed to cool slowly to about 25°C and charged in a suitable vessel to obtain ointment for external use.

(Example 12) Ointment for external use

5 Prescription:

The ointment is composed of tinidazole (2 g), propylene glycol (10 g), polyoxyethylene glycol monostearate (5 g), liquid paraffin (20 g), white petrolatum (60 g), and distilled water (an amount making total 100 g).

10 Preparation method:

Distilled water and propylene glycol were mixed and a temperature of the mixture was adjusted to 70°C, followed by stirring. To the above mixture were added polyoxyethylene glycol monostearate, tinidazole, liquid paraffin and white petrolatum mixed at a temperature of 70°C. The mixture was allowed to cool slowly to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel to obtain ointment for external use.

20 (Example 13) Cream for external use

Prescription:

The cream is composed of tinidazole (1.8 g), stearic acid (5 g), polyoxyethylene cetostearyl ether (12E.O.) (0.5 g), polyoxyethylene cetostearyl ether (20E.O.) (0.5 g), cetanol (5 g), cetyl octanoate (5 g), liquid paraffin (5 g), beeswax (1 g), glycerin (5 g), 1,3-butylene glycol (5 g), triethanolamine (5 g), hydrochloric acid (2.7 g), distilled water (an amount making total 100 g).

Preparation method:

As oil phase, stearic acid, polyoxyethylene cetostearyl ether (12E.O.), polyoxyethylene cetostearyl ether (20E.O.), cetanol, cetyl octanoate, liquid paraffin, and beeswax were melted and stirred at a temperature of about 70 to 75°C. To the above mixture were slowly added, while keeping a temperature at about 70°C, a solution of distilled water, glycerin, 1,3-butylene glycol and triethanolamine dissolved therein. Then, a dissolved solution of distilled water,

tinidazole and hydrochloric acid was heated to about 70°C and slowly added to the above mixture. The formed emulsified liquid was continuously stirred and after cooling it at a temperature of about 25°C, then, charged in a suitable vessel to obtain cream.

(Example 14) Cream for external use

Prescription:

The cream is composed of tinidazole (1.8 g), stearic acid (3 g), glycol monostearate (4 g), polyoxyethylene glycol monostearate (1 g), polyoxyethylene cetostearyl ether (12E.O.) (0.5 g), polyoxyethylene cetostearyl ether (20E.O.) (0.5 g), cetanol (5 g), liquid paraffin (10 g), cetyl octanoate (5 g), ethyl paraoxybenzoate (0.3 g), silicone oil (1 g), beeswax (1.5 g), 1,3-butylene glycol (7 g), glycerin (5 g), sodium hydroxide (suitable amount), hydrochloric acid (suitable amount), and distilled water (an amount making total 100 g).

Preparation method:

To a dissolved solution of distilled water, 1,3-butylene glycol and glycerin, tinidazole was added and further hydrochloric acid was added thereto until tinidazole was completely dissolved. This liquid was heated to a temperature of about 70°C and a pH thereof was made 6.9 with sodium hydroxide. This mixture was slowly added while stirring to a molten liquid in which stearic acid, glycol monostearate, polyoxyethylene glycol monostearate, polyoxyethylene cetostearyl ether (12E.O.), polyoxyethylene cetostearyl ether (20E.O.), cetanol, liquid paraffin, cetyl octanoate, ethyl paraoxybenzoate, silicone oil and beeswax were mixed and adjusted to a temperature of about 70 to 75°C. The formed emulsion was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel to obtain cream.

(Example 15) Cream for external use

Prescription:

The cream is composed of tinidazole (2 g), n-octadecyl alcohol (5 g), stearic acid (5 g), triethanolamine (5 g), liquid paraffin (8 g), disodium edetate (0.2 g), glycerin (10 g), thymol (0.2 g), hydrochloric acid (suitable amount),  
5 distilled water (an amount making total 100 g).

Preparation method:

A mixture of n-octadecyl alcohol, stearic acid and liquid paraffin was melted by heating while stirring and a temperature was adjusted to about 70°C, and tinidazole was  
10 then added thereto. To the resulting mixture was slowly added a dissolved material of distilled water, glycerin and triethanolamine adjusted a temperature to about 70°C while stirring. After adjusting a pH thereof to 6.8 by the  
15 addition of hydrochloric acid, disodium edetate and thymol were added thereto while continuously stirring. The mixture was cooled to a temperature of about 25°C and then charged in a suitable vessel to obtain cream for external use.

(Example 16) Lotion for external use

20 Prescription:

The lotion is composed of tinidazole (2 g), stearic acid (3 g), cetanol (1 g), polyoxyethylene cetostearyl ether (20E.O.) (0.5 g), triethanolamine (0.2 g), glycerin (5 g), isopropanol (10 g), and distilled water (an amount making  
25 total 100 g).

Preparation method:

Cetanol, polyoxyethylene cetostearyl ether (20E.O.), stearic acid and tinidazole were melted by heating while stirring, and a molten mixture of triethanolamine, distilled  
30 water and glycerin was further added thereto. Next, the resulting mixture was cooled to a temperature of 40°C, isopropanol was added thereto and the mixture was rapidly cooled to a temperature of about 25°C while continuously stirring. After cooling, the mixture was charged in a  
35 suitable vessel to obtain lotion for external use.

(Example 17) Lotion for external use

Prescription:

The lotion is composed of tinidazole (1.8 g),  
isopropanol (10 g), n-octadecyl alcohol (10 g), cetanol (5 g),  
Tween 80 (2 g), 1,3-butylene glycol (10 g), sodium  
5 carboxymethyl cellulose (3 g), and distilled water (an amount  
making total 100 g).

Preparation method:

n-Octadecyl alcohol and cetanol were melted by heating  
and slowly added to a heated mixture of distilled water,  
10 sodium carboxymethyl cellulose, Tween 80, 1,3-butylene glycol  
and tinidazole. Then, the resulting mixture was cooled to a  
temperature of about 40°C and isopropanol was added thereto,  
and the mixture was cooled to about 25°C while continuously  
stirring. The resulting material was charged in a suitable  
15 vessel to obtain lotion for external use.

(Example 18) Moisture patch

Prescription:

The patch is composed of tinidazole (2 g), kaolin (5 g),  
20 liquid paraffin (10 g), glycerin (15 g), sodium carboxymethyl  
cellulose (5 g), crotamiton (1.5 g), zinc oxide (2 g), Tween  
80 (2 g), gelatin (5 g), sodium polyacrylate (5 g), and  
distilled water (an amount making total 100 g).

Preparation method:

25 To distilled water were added a material in which  
sodium carboxymethyl cellulose and gelatin were melted by  
heating and added, and then, kaolin was added to disperse  
therein. This material was added under stirring to a  
material in which zinc oxide, sodium polyacrylate and liquid  
30 paraffin had been dispersed by stirring. Further, to the  
above mixture was added a material in which tinidazole,  
crotamiton, glycerin and Tween 80 had been stirred and heated,  
and a temperature of which had been adjusted to about 60°C  
under stirring and heating. Ointment thus obtained was  
35 applied with 1000g/m<sup>2</sup> to an unwoven fabric and the fabric was  
cut into a size of 10 cm x 14 cm (280 mg of tinidazole was  
contained per 14 g of ointment) to obtain a patch.

(Example 19) Moisture patch

Prescription:

The patch is composed of tinidazole (2 g), sorbitan  
5 monooleate (0.5 g), polyoxyethylene sorbitan monooleate (0.5  
g), castor oil (1 g), crotamiton (1 g), gelatin (1 g), kaolin  
(12 g), sodium metaphosphate (0.15 g), 1,3-butylene glycol (5  
g), starch grafted acrylate 300 (2 g), sodium polyacrylate (5  
g), methacrylic acid-n-butyl acrylate copolymer (4 g), D-  
10 sorbitol solution (70%) (50 g), tartaric acid (1.7 g),  
titanium oxide (1 g), magnesium hydroxide-aluminum hydroxide  
co-precipitate (0.25 g), dibutylhydroxytoluene (0.2 g), and  
distilled water (an amount making total 100 g).

Preparation method:

15 Suitable amounts of distilled water and D-sorbitol  
solution were mixed and dissolved. To this mixture was added  
titanium oxide, and then, suitable amounts of kaolin and a D-  
sorbitol solution were added to the mixture under stirring.  
To this mixture was added gelatin, and then, a methacrylic  
20 acid-n-butyl acrylate copolymer. To this mixture was further  
added a material obtained by mixing while stirring a mixed  
material in which sodium polyacrylate, starch grafted  
acrylate 300, magnesium hydroxide-aluminum hydroxide co-  
precipitate, 1,3-butylene glycol and castor oil were  
25 dissolved, a material in which tinidazole, crotamiton and  
dibutylhydroxytoluene were dispersed under heating, and a  
mixture of sorbitan monooleate and polyoxyethylene sorbitan  
monooleate. Then, to the resulting mixture was added a  
material in which sodium metaphosphate was dissolved in a  
30 small amount of distilled water, and finally, was added a  
mixture of a remaining D-sorbitol solution and tartaric acid  
adjusted to a temperature of about 60°C under stirring.  
Ointment thus obtained was applied with 1000 g/m<sup>2</sup> to an  
unwoven fabric and the fabric was cut into a size of 10 cm x  
35 14 cm (280 mg of tinidazole was contained per 14 g of  
ointment) to obtain a patch.

(Example 20) Dry patch (plaster)

Prescription:

The patch is composed of tinidazole (2 g), liquid paraffin (8 g), dibutylhydroxytoluene (0.2 g), crotamiton (1 g), polyoxyethylene glycol monostearate (2 g), polyoxyethylene cetostearyl ether (20E.O.) (1.8 g), methacrylic acid-n-butyl acrylate copolymer (5 g), myristyl alcohol (8 g), natural rubber latex (as a solid material) (20 g), synthetic rubber SBR latex (as a solid material) (37 g), and polybutene (15 g).

Preparation method:

Tinidazole, dibutylhydroxytoluene and crotamiton were dispersed by stirring under heating. Polyoxyethylene glycol monostearate, polyoxyethylene cetostearyl ether (20E.O.) and myristyl alcohol were added to the above mixture and the resulting mixture was mixed under heating. The resulting mixture was added to a molten state mixture of a methacrylic acid-n-butyl acrylate copolymer, a natural rubber latex and a synthetic rubber SBR latex while continuously stirring. To this mixture were also further added liquid paraffin and polybutene while continuously stirring. Ointment thus obtained was applied with 100 g/m<sup>2</sup> to an unwoven or woven fabric. After drying, the fabric was cut into a size of 10 cm x 14 cm (28 mg of tinidazole was contained per 1.4 g of ointment) to obtain plaster.

(Example 21) Cream for external use

Prescription:

The cream is composed of metronidazole (2 g), clotrimazole (0.1 g), clobetasol propionate (0.005 g), glycol monostearate (10 g), cetanol (7 g), liquid paraffin (9 g), white petrolatum (3.5 g), propylene glycol (6.5 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g).

Preparation method:

Glycol monostearate, cetanol, liquid paraffin and white petrolatum were stirred under heating to about 85°C, and to

the mixture was added a mixture of propylene glycol, sodium lauryl sulfate and purified water prepared by stirring under heating to about 85°C. Then, under stirring, metronidazole, clotrimazole and clobetasol propionate were added thereto.

- 5 The resulting mixture was cooled to about 25°C while continuously stirring and the resulting cream was charged in a suitable vessel.

(Example 22) Cream for external use

10 Prescription:

The cream is composed of (a) metronidazole (2 g), lidocaine (0.05 g), prednisolone valerate acetate (0.005 g); (b) glycol monostearate (10 g), cetanol (7 g), liquid paraffin (9 g), white petrolatum (3.5 g); and (c) propylene glycol (6.5 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g).

Preparation method:

- 20 Component(b) was stirred under heating at about 85°C, and to the mixture was added component(c) which had been stirred under heating to about 85°C, and component(a) was added to the above mixture while stirring. The mixture was cooled to about 25°C while continuously stirring, and the resulting cream was charged in a suitable vessel.

25 (Example 23) Cream for external use

Prescription:

- 30 The cream is composed of tinidazole (2 g), clotrimazole (0.1 g), clobetasol propionate (0.002 g), glycol monostearate (10 g), cetanol (7 g), liquid paraffin (9 g), white petrolatum (3.5 g), propylene glycol (6.5 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g).

Preparation method:

- 35 Glycol monostearate, cetanol, liquid paraffin and white petrolatum were stirred under heating to about 85°C, and to the mixture was added a mixture of propylene glycol, sodium lauryl sulfate and purified water prepared by stirring under

heating to about 85°C. Then, under stirring, to the mixture were added tinidazole, clotrimazole and clobetasol propionate. The mixture was cooled to about 25°C while continuously stirring, and the resulting cream was charged in a suitable vessel.

(Example 24) Cream for external use

Prescription:

The cream is composed of tinidazole (2 g), chloramphenicol (0.001 g), hydrocortisone acetate (0.001 g), glycol monostearate (8 g), cetanol (7 g), liquid paraffin (10 g), white petrolatum (3.5 g), propylene glycol (6.5 g), sodium lauryl sulfate (0.8 g), and purified water (an amount making total 100 g).

Preparation method:

Glycol monostearate, cetanol, liquid paraffin and white petrolatum were stirred under heating to about 85°C, and to the mixture were added a mixture of propylene glycol, sodium lauryl sulfate and purified water prepared by stirring under heating to about 85°C. Then, under stirring, to the mixture were added tinidazole, chloramphenicol and hydrocortisone acetate. The mixture was cooled to about 25°C while continuously stirring, and the resulting cream was charged in a suitable vessel.

(Example 25) Gel

Prescription:

The gel is composed of (a) tinidazole (3 g), diphenhydramine hydrochloride (0.2 g), betamethasone (0.01 g), carpronium chloride (0.2 g); (b) polyoxyethylene oleyl alcohol ether (1 g); (c) polyoxyethylene glycol 1500 (6 g), polyoxyethylene glycol 400 (2 g), disodium EDTA (0.2 g); (d) dipropylene glycol (8 g); (e) aqueous phase: potassium hydroxide (0.1 g); (f) carboxyvinyl polymer (0.5 g), methyl cellulose (0.2 g), and purified water (an amount making total 100 g).

Preparation method:

After (f) was homogeneously dissolved, (c) was added thereto and (a) was added thereto, and the mixture was heated, dissolved and dispersed. Subsequently, (b) was added to (d) and the mixture was melted by heating at about 60°C, and added to the above dispersion. The mixture was neutralized by the addition of (e) while stirring and cooled to a temperature of about 25°C. The resulting gel was charged in a suitable vessel.

10 (Example 26) Cream for external use

Prescription:

The cream is composed of metronidazole (2 g), glycol monostearate (10.4 g), cetanol (7.3 g), liquid paraffin (9 g), propylparaben (0.05 g), white petrolatum (3.5 g), propylene glycol (6.5 g), sodium lauryl sulfate (1 g), methylparaben (0.05 g), urea (2 g), and purified water (an amount making total 100 g).

Preparation method:

Glycol monostearate, cetanol, liquid paraffin, propylparaben and white petrolatum were stirred while heating at about 85°C. A mixture of propylene glycol, sodium lauryl sulfate, methylparaben, urea and purified water were stirred while heating at about 85°C and added thereto. Metronidazole was added thereto while stirring. The mixture was cooled to about 25°C while continuously stirring and then charged in a suitable vessel.

(Example 27) Shampoo

Prescription:

The shampoo is composed of metronidazole (2 g), polyglycerin monolaurate (4 g), polyoxyethylene lauryl ether sodium sulfate (7 g), lauryl dimethylaminoacetate betaine (2.5 g), coconut fatty acid diethanol amide (4 g), polyethylene glycol (5 g), 1,3-butylene glycol (3 g), citric acid (suitable amount), and purified water (to make total amount 100 g).

Preparation method:

Metronidazole is added to a mixture containing suitable amounts of polyethylene glycol and purified water and the mixture is melted by heating. In other vessel are weighed suitable amounts of polyglycerin monolaurate, polyoxyethylene lauryl ether sodium sulfate, lauryl dimethylacetate betaine, coconut fatty acid diethanol amide, polyethylene glycol, 1,3-butylene glycol and purified water, and the resulting mixture is heated to about 70°C while stirring and added to a mixture of metronidazole, polyethylene glycol and purified water. The mixture is adjusted to pH about 6.5 with citric acid, and is cooled until a temperature becomes about 25°C under stirring.

(Example 28) Gel

Prescription:

The gel is composed of metronidazole (1 g), polyethylene glycol (8 g), carboxyvinyl polymer (0.5 g), methyl cellulose (0.2 g), propylene glycol (5 g), glycerin (2 g), polyoxyethylene oleyl cetyl ether (1 g), isopropanol (5 g), sodium hydroxide (suitable amount), and purified water (an amount making total 100 g)

Preparation method:

Polyethylene glycol was added to purified water and dissolved, and metronidazole was further added thereto and dissolved by heating. The solution was cooled to about 50°C, and to the solution was added a material in which polyoxyethylene cetyl ether was added to propylene glycol and glycerin and heated to about 50°C under stirring. While further continuously stirring, sodium hydroxide was added thereto and a pH thereof was adjusted to about 6.8. The mixture was further cooled to about 40°C, isopropanol was then added thereto, and after cooling the mixture to about 25°C, it was charged in a suitable vessel.

(Example 29) Ointment

Prescription:

The ointment is composed of (a) metronidazole (2 g),

crotamiton (2 g); (b) stearic acid (2 g), glycol monostearate (12 g), polyoxyethylene glycol monostearate (3 g), polyoxyethylene cetostearyl ether (12E.O.) (1 g), polyoxyethylene cetostearyl ether (20E.O.) (1 g), cetanol (2 g), liquid paraffin (8 g); and (c) 1,3-butylene glycol (7 g), glycerin (5 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 75°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 75°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 30) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (2 g), lidocaine (0.05 g), prednisolone valerate acetate (0.005 g); (b) glycol monostearate (10 g), cetanol (7 g), liquid paraffin (9 g), white petrolatum (3.5 g); (c) propylene glycol (6.5 g), sodium lauryl sulfate (1 g), purified water (an amount making the total amount 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 31) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (2.5 g), ketoconazole (0.1 g); (b) stearic acid (5 g), stearyl alcohol (5 g), liquid paraffin (5 g), isopropyl myristate (1 g), Span 60 (1 g), thymol(0.2 g); and (c) Tween 60 (0.5 g), propylene

glycol (5 g), triethanolamine (0.4 g), and distilled water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C.

5 To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

10

(Example 32) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (3 g), pipemidic acid trihydrate (0.1 g), prednisolone (0.001 g);  
15 (b) glycol monostearate (5 g), polyoxyethylene (23) cetyl ether (2 g), cetanol (6 g), white petrolatum (5 g), liquid paraffin (5 g), caprylic/capric acid triglyceride (5 g), octyl dodecyl myristate (3 g), propyl parahydroxybenzoate (0.1 g); and (c) propylene glycol (5 g), methyl  
20 parahydroxybenzoate (0.1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C.

To the solution was added under stirring a material in which  
25 component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

30 (Example 33) Ointment for external use

Prescription:

The ointment is composed of (a) metronidazole (2 g), crotamiton (1 g), fluorocinolone acetonide (0.001 g); (b)  
white petrolatum (45 g), cetanol (20 g), polyoxyethylene  
35 hydrogenated castor oil (5 g), Tween 80 (2 g), liquid paraffin (5 g), propyl parahydroxybenzoate(0.1 g); and (c) methyl parahydroxybenzoate (0.1 g), and distilled water (an

amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

10 (Example 34) Ointment for external use

Prescription:

The ointment is composed of (a) metronidazole (2 g), diphenhydramine hydrochloride (0.2 g), lidocaine (0.1 g); (b) stearyl alcohol (7 g), cetanol (3 g), white petrolatum (30 g), glycol monostearate (10 g), span 80 (1.5 g), liquid paraffin (5 g); and (c) propylene glycol (5 g), Tween 80 (1 g), and distilled water (an amount making total 100 g)

Preparation method:

Component(c) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(b) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

25

(Example 35) Ointment for external use

Prescription:

The ointment is composed of (a) metronidazole (2 g), gentamicin sulfate (0.005 g); (b) glycol monostearate (15 g), polyoxyethylene glycol monostearate (3 g), polyoxyethylene cetostearyl ether (2 g), cetanol (5 g), beeswax (5 g), white petrolatum (20 g), and (c) distilled water (an amount making the total amount 100 g)

Preparation method:

35 Component(c) was dissolved and adjusted to about 85°C. To the solution was added a material in which component(b)

had been dissolved and adjusted to about 85°C under stirring, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

5

(Example 36) Lotion

Prescription:

The lotion is composed of (a) metronidazole (2 g),  
betamethasone valerate (0.005 g), bifonazole (0.05 g); (b)  
10 stearic acid (2 g), cetanol (1.5 g), petrolatum (4 g),  
squalane (5 g), caprylic/capric acid triglyceride (2 g),  
sorbitan monooleate (2 g), polyethylene glycol (5 g); (c)  
dipropylene glycol (5 g), triethanol amine (0.7 g), purified  
water (60 g); (d) isopropanol (10 g), and purified water (an  
15 amount making total 100 g)

Preparation method:

Component(c) was dissolved and adjusted to about 75°C.  
To the solution was added under stirring a material in which  
component(b) had been dissolved and adjusted to about 75°C,  
20 followed by the addition of component(a). The mixture was  
rapidly cooled to a temperature of about 40°C while  
continuously stirring, then component(d) was added thereto,  
and the mixture was cooled to about 25°C under stirring. The  
resulting lotion was charged in a suitable sealed vessel.

25

(Example 37) Patch

Prescription:

The patch is composed of (a) metronidazole (3 g),  
crotamiton (1 g), prednisolone (0.05 g); (b) D-sorbitol  
30 (70%) (30 g), purified water (9 g), kaolin (13 g), titanium  
oxide (1 g); (c) gelatin (1 g), purified water (4 g); (d)  
sodium metaphosphate (0.1 g), purified water (1 g); (e)  
sodium polyacrylate (5 g), starch grafted acrylate 300 (1 g),  
propylene glycol (5 g), castor oil (1 g), magnesium  
35 hydroxide-aluminum hydroxide co-precipitate (0.25 g),  
sorbitan monooleate (0.5 g), polyoxyethylene sorbitan  
monooleate (0.5 g); (f) D-sorbitol (70%) (14 g),

dibutylhydroxytoluene (0.2 g); (g) methacrylic acid and n-butyl acrylate copolymer (3 g); and (h) D-sorbitol (70%) (4.9 g), tartaric acid (1.5 g)

Preparation method:

5           Component(b) was dissolved and adjusted to about 40°C. To the solution was added under stirring a material in which component(d) had been dissolved and adjusted to about 60°C, followed by the addition of component(c) and then component(g). To the resulting mixture was added a material  
10 in which components(a) and (e) had been well mixed, followed by the addition of component(f), and then, component(h) was added thereto little by little under stirring. 14 g of ointment thus obtained was applied evenly to an unwoven fabric having a size of 10 cm x 14 cm to obtain a patch.

15           (Example 38) Patch (Plaster)

Prescription:

The plaster is composed of (a) metronidazole(3 g), indometacin (1 g); (b) liquid paraffin (7 g), isopropyl  
20 myristate (3 g), polybutene (15 g), 1,3-pentadiene copolymer resin (26 g); (c) polyoxyethylene sorbitan monostearate (1.5 g), zinc oxide (3 g), titanium oxide (2 g), dibutylhydroxytoluene (0.2 g), crotamiton (1 g); (d) kaolin (6 g); (e) natural rubber latex (as a solid material) (15 g),  
25 synthetic rubber SBR (as a solid material) (17 g); and (f) glycerin (0.25 g), purified water (1 g), and sodium polyacrylate (0.05 g)

Preparation method:

          component(b) was mixed and melted at about 110°C and a  
30 temperature thereof was adjusted to about 90°C. To this mixture was added component(a), and after adjusting to about 70°C, a material in which component(c) and (d) had been mixed was added to the above mixture. Under further stirring, component(f) was added and component(e) was also added  
35 thereto at about 70°C. 14 g of ointment thus obtained was applied evenly to an unwoven fabric in a size of 10 cm x 14 cm to obtain a patch.

(Example 39) Gel

Prescription:

The gel is composed of (a) metronidazole (3 g),  
5 diphenhydramine hydrochloride (0.5 g), betamethasone (0.01  
g); (b) polyoxyethylene oleyl alcohol ether (1 g); (c)  
polyethylene glycol 1500 (6 g), polyoxyethylene glycol 400 (2  
g), disodium EDTA (0.2 g); (d) dipropylene glycol (8 g); (e)  
potassium hydroxide (0.1 g); (f) carboxyvinyl polymer (0.5 g),  
10 methylcellulose (0.2 g), purified water (an amount making  
total 100 g)

Preparation method:

After component(f) was homogeneously dissolved,  
component(c) was added to the solution, and then,  
15 component(a) was added to the same to dissolve or disperse  
therein under heating. To the resulting material was added a  
material in which component(b) had been added to component(d)  
and mixed to melt at about 60°C. Moreover, under stirring,  
component(e) was added thereto while stirring to neutralize  
20 the mixture, and the mixture was cooled to about 25°C. The  
resulting gel was charged in a suitable vessel.

(Example 40) Cream for external use

Prescription :

25 The cream is composed of (a) tinidazole (1 g),  
prednisolone valerate acetate (0.005 g); (b) glycol  
monostearate (8 g), cetanol (7 g), liquid paraffin (10 g),  
white petrolatum (3.5 g); (c) propylene glycol (6.5 g),  
sodium lauryl sulfate (1 g), purified water (an amount making  
30 total 100 g).

Preparation method:

Component(b) was dissolved and adjusted to about 85°C.  
To the solution was added under stirring a material in which  
component(c) had been dissolved and adjusted to about 85°C,  
35 followed by the addition of component(a). The mixture was  
cooled to a temperature of about 25°C while continuously  
stirring, and then, charged in a suitable vessel.

(Example 41) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (2.5 g),  
5 azelastine hydrochloride (0.02 g), prednisolone acetate  
(0.001g); (b) glycol monostearate(5 g), polyoxyethylene(23)  
cetyl ether (2 g), cetanol(5 g), white petrolatum (3.5 g),  
liquid paraffin (5 g), isopropyl myristate (5 g), octyl  
dodecyl myristate (3 g), propyl parahydroxybenzoate (0.15 g);  
10 (c) propylene glycol (7 g), methyl parahydroxybenzoate (0.15  
g), distilled water (an amount making the total amount 100 g).

Preparation method:

Component(b) was dissolved and adjusted to about 85°C.  
To the solution was added under stirring a material in which  
15 component(c) had been dissolved and adjusted to about 85°C,  
followed by the addition of component(a). The mixture was  
cooled to a temperature of about 25°C while continuously  
stirring, and then, charged in a suitable vessel.

20 (Example 42) Cream

Prescription:

The cream is composed of (a) tinidazole (2.0 g),  
tolnaftate (0.05 g), (b) stearic acid (5 g), stearyl  
alcohol(5 g), liquid paraffin (5 g), isopropyl myristate (1  
25 g), Span 60 (1.2 g), thymol(0.2 g); and (c) Tween 60 (0.7 g),  
propylene glycol (6 g), triethanolamine (0.4 g), and purified  
water (an amount making the total amount 100 g).

Preparation method:

Component(b) was dissolved and adjusted to about 85°C.  
30 To the solution was added under stirring a material in which  
component(c) had been dissolved and adjusted to about 85°C,  
followed by the addition of component(a). The mixture was  
cooled to a temperature of about 25°C while continuously  
stirring, and then, charged in a suitable vessel.

35

(Example 43) Cream

Prescription:

The cream is composed of (a) tinidazole (2 g),  
aciclovir (0.2 g); (b) stearic acid (5 g), stearyl alcohol (5  
g), liquid paraffin (5 g), isopropyl myristate (1 g), Span 60  
(1.2 g), thymol(0.2 g); (c) Tween 60 (0.7 g), propylene  
5 glycol (6 g), triethanolamine (0.4 g), purified water (an  
amount making total 100 g).

Preparation method:

Component(b) was dissolved and adjusted to about 85°C.  
To the solution was added under stirring a material in which  
10 component(c) had been dissolved and adjusted to about 85°C,  
followed by the addition of component(a). The mixture was  
cooled to a temperature of about 25°C while continuously  
stirring, and then, charged in a suitable vessel.

15 (Example 44) Ointment for external use

Prescription:

The ointment is composed of (a) tinidazole (2 g),  
diclofenac sodium (0.05 g), crotamiton (1 g), fluocinolone  
acetoneide (0.001 g); (b) white petrolatum (45 g), cetanol (20  
20 g), polyoxyethylene hydrogenated castor oil (5 g), Tween 80  
(2 g), liquid paraffin (5 g), propyl parahydroxybenzoate (0.1  
g); and (c) methyl parahydroxybenzoate (0.1 g), and purified  
water (an amount making the total amount 100 g).

Preparation method:

25 Component(b) was dissolved and adjusted to about 85°C.  
To the solution was added under stirring a material in which  
component(c) had been dissolved and adjusted to about 85°C,  
followed by the addition of component(a). The mixture was  
cooled to a temperature of about 25°C while continuously  
30 stirring, and then, charged in a suitable vessel.

(Example 45) Ointment for external use

Prescription:

The ointment is composed of (a) tinidazole (2 g),  
35 extract of calves blood (1 g), diphenhydramine hydrochloride  
(0.2 g), lidocaine (0.1 g); (b) stearyl alcohol (7 g),  
cetanol (3 g), white petrolatum (30 g), glycol monostearate

(10 g), span 80 (1.5 g), liquid paraffin (5 g); and (c) propylene glycol (5 g), Tween 80 (1 g), and distilled water (an amount making the total amount 100 g).

Preparation method:

5       Component(c) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(b) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously  
10       stirring, and then, charged in a suitable vessel.

(Example 46) Ointment for external use

Prescription:

15       The ointment is composed of (a) tinidazole (2 g), gentamicin sulfate (0.005 g), (b) glycol monostearate (15 g), polyoxyethylene glycol monostearate (3 g), polyoxyethylene cetostearyl ether (2 g), cetanol (5 g), beeswax (5 g), white petrolatum (20 g); (c) distilled water (an amount making the total amount 100 g).

20       Preparation method:

      Component(c) was adjusted to about 85°C. To the solution was added under stirring a material in which component(b) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was  
25       cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 47) Lotion

Prescription:

30       The lotion is composed of (a) tinidazole (2 g), ofloxacin (0.005 g), clotrimazole (0.05 g); (b) stearic acid (2 g), cetanol (1.5 g), white petrolatum (4 g), squalane (5 g), caprylic/caproic acid triglyceride (2 g), sorbitan monooleate (2 g), polyethylene glycol (5 g); (c) dipropylene  
35       glycol (5 g), triethanolamine (0.7 g), purified water (60 g); and (d) isopropanol (10 g), purified water (an amount making total 100 g).

Preparation method:

Component(c) was dissolved and adjusted to about 70°C. To the solution was added under stirring a material in which component(b) had been dissolved and adjusted to about 70°C, followed by the addition of component(a). The mixture was rapidly cooled to a temperature of about 40°C while continuously stirring, then component(d) was added thereto, and the mixture was cooled to about 25°C under stirring. The resulting lotion was charged in a suitable sealed vessel.

(Example 48) Patch

Prescription:

The patch is composed of (a) tinidazole (3 g), crotamiton (1 g), prednisolone (0.05 g); (b) D-sorbitol (70%) (30 g), purified water (9 g), kaolin (13 g), titanium oxide (1 g); (c) gelatin (1 g), purified water (4 g); (d) sodium metaphosphate (0.1 g), purified water (1 g); (e) sodium polyacrylate (5 g), starch grafted acrylate 300 (1 g), propylene glycol (5 g), castor oil (1 g), magnesium hydroxide-aluminum hydroxide co-precipitate (0.25 g), sorbitan monooleate (0.5 g), sorbitan polyoxyethylene monooleate (0.5 g); (f) D-sorbitol (70%) (14 g), dibutylhydroxyltoluene (0.2 g); (g) methacrylic acid-n-butyl acrylate copolymer (3 g); and (h) D-sorbitol (70%) (4.9 g), tartaric acid (1.5 g).

Preparation method:

A temperature of component(b) was adjusted to about 40°C, and a material in which a temperature of component(d) had been adjusted to about 60°C was added to component(b) under stirring. Then, component(c) was added to the above mixture and component(g) was added thereto while stirring. To this mixture was added a material in which component(a) and component(e) had been well mixed, followed by the addition of component(f), and component(h) is added thereto while stirring. From the resulting ointment, 14 g was weighed and applied evenly to an unwoven fabric in a size of 10 cm x 14 cm to obtain a patch.

(Example 49) Patch (plaster)

Prescription:

The patch is composed of (a) tinidazole (3 g),  
5 indometacin (1 g); (b) liquid paraffin (7 g), isopropyl  
myristate (3 g), polybutene (15 g), 1,3-pentadiene copolymer  
resin (26 g); (c) polyoxyethylene sorbitan monostearate (1.5  
g), zinc oxide (3 g), titanium oxide (2 g),  
dibutylhydroxyltoluene (0.2 g), crotamiton (1 g); (d) kaolin  
10 (6 g); (e) natural rubber latex (as a solid material) (15 g),  
synthetic rubber SBR (as a solid material) (17 g); (f)  
glycerin (0.25 g), purified water (1 g), sodium polyacrylate  
(0.05 g).

Preparation method:

15 Component(b) was mixed and melted at a temperature of  
about 110°C, and then, adjusted to about 90°C, and  
component(a) was added thereto and the resulting mixture was  
adjusted to a temperature of about 70°C. To the mixture was  
added a material in which mixture of component(c) and (d) had  
20 been mixed. Further, component(f) was added thereto and (e)  
was added at a temperature of about 70°C. The resulting  
ointment was applied with 100 g/m<sup>2</sup> to an unwoven or woven  
fabric and the fabric was cut into a size of 10 cm × 14 cm.

25 (Example 50) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (1 g),  
fluorouracil (0.02 g), prednisolone valerate acetate (0.005  
g); (b) glycol monostearate (8 g), cetanol (7 g), liquid  
30 paraffin (10 g), white petrolatum (3.5 g); and (c) propylene  
glycol (6.5 g), sodium lauryl sulfate (1 g), and purified  
water (an amount making the total amount 100 g).

Preparation method:

Component(b) was dissolved and adjusted to about 85°C.  
35 To the solution was added under stirring a material in which  
component(c) had been dissolved and adjusted to about 85°C,  
followed by the addition of component(a). The mixture was

cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 51) Cream for external use

5 Prescription:

The cream is composed of metronidazole (2 g), glycol monostearate (10 g), cetanol (7 g), liquid paraffin (9 g), white petrolatum (3.5 g), propylene glycol (6.5 g), sodium lauryl sulfate (1 g), and purified water (an amount making  
10 the total amount 100 g)

Preparation method:

Glycol monostearate, cetanol, liquid paraffin and white petrolatum were mixed by stirring under heating at about 85°C. To the above mixture was added a material in which propylene  
15 glycol, sodium lauryl sulfate and purified water had been mixed by stirring under heating at about 85°C, and metronidazole was added thereto while stirring. The resulting cream was charged in a suitable vessel.

20 (Example 52) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (0.5 g); (b) glycol monostearate (10 g), cetanol (7 g), liquid paraffin (9 g), white petrolatum (3.5 g); and (c) propylene glycol (6.5  
25 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which  
30 component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

35 (Example 53) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (0.5 g); (b)

glycol monostearate (10 g), cetanol (7 g), liquid paraffin (9 g), white petrolatum (3.5 g); (c) propylene glycol (6.5 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

5 Preparation method:

Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 54) Cream for external use

Prescription:

15 The cream is composed of (a) metronidazole (1.5 g), ketoconazole (0.1 g); (b) glycol monostearate(5 g), polyoxyethylene (23) cetyl ether (2 g), stearic acid (0.5 g), cetanol (5 g), white petrolatum (3.5 g), liquid paraffin (5 g), isopropyl myristate (5 g), octyl dodecyl myristate (3 g),  
20 propyl parahydroxybenzoate (0.15 g); and (c) propylene glycol (7 g), methyl parahydroxybenzoate (0.15 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

30

(Example 55) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (1.5 g), ketoconazole (0.1 g); (b) glycol monostearate(5 g),  
35 polyoxyethylene (23) cetyl ether (2 g), stearic acid (0.5 g), cetanol (5 g), white petrolatum (3.5 g), liquid paraffin (5 g), isopropyl myristate (5 g), octyl dodecyl myristate (3 g),

propyl parahydroxybenzoate (0.15 g); and (c) propylene glycol (7 g), methyl parahydroxybenzoate (0.15 g), and distilled water (an amount making total 100 g)

Preparation method:

5           Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously  
10 stirring, and then, charged in a suitable vessel.

(Example 56) Cream for external use

Prescription:

15           The cream is composed of (a) metronidazole (3 g), norfloxacin (0.2 g); (b) stearic acid (5 g), stearyl alcohol (5 g), liquid paraffin (5 g), isopropyl myristate (1 g), Span 60 (1.2 g), thymol (0.2 g); and (c) Tween 60 (0.7 g), propylene glycol (6 g), triethanolamine (0.4 g), and purified water (an amount making total 100 g)

20           Preparation method:

            Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was  
25 cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 57) Cream for external use

Prescription:

30           The cream is composed of (a) tinidazole (3 g), norfloxacin (0.2 g); (b) stearic acid (5 g), stearyl alcohol (5 g), liquid paraffin (5 g), isopropyl myristate (1 g), Span 60 (1.2 g), thymol (0.2 g); and (c) Tween 60 (0.7 g), propylene glycol (6 g), triethanolamine (0.4 g), and purified  
35 water (an amount making total 100 g)

Preparation method:

            Component(b) was dissolved and adjusted to about 85°C.

To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 58) Ointment for external use

Prescription:

The ointment is composed of (a) metronidazole (2 g), diclofenac sodium (0.1 g); (b) white petrolatum (45 g), cetanol (20 g), polyoxyethylene hydrogenated castor oil (5 g), Tween 80 (2 g), crotamiton (3 g), liquid paraffin (5 g), propyl parahydroxybenzoate (0.1 g); and (c) methyl parahydroxybenzoate (0.1 g), distilled water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 59) Ointment for external use

Prescription:

The ointment is composed of (a) tinidazole (2 g), diclofenac sodium (0.1 g); (b) white petrolatum (45 g), cetanol (20 g), polyoxyethylene hydrogenated castor oil (5 g), Tween 80 (2 g), crotamiton (3 g), liquid paraffin (5 g), propyl parahydroxybenzoate (0.1 g); and (c) methyl parahydroxybenzoate (0.1 g), and distilled water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was

cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 60) Cream for external use

5 Prescription:

The cream is composed of (a) metronidazole (10 g); (b) glycol monostearate (7 g), polyoxyethylene glycol monostearate (3 g), polyoxyethylene cetostearyl ether (2 g), polyoxyethylene hydrogenated castor oil (1 g), cetanol (5 g),  
10 beeswax (1 g), liquid paraffin (3 g); and (c) polyethylene glycol (5 g), 1,3-butylene glycol (4 g), and distilled water (an amount making total 100 g)

Preparation method:

Component(c) was dissolved and adjusted to about 85°C.  
15 To the solution was added under stirring a material in which component(b) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

20

(Example 61) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (10 g); (b) glycol monostearate (7 g), polyoxyethylene glycol  
25 monostearate (3 g), polyoxyethylene cetostearyl ether (2 g), polyoxyethylene hydrogenated castor oil (1 g), cetanol (5 g), beeswax (1 g), liquid paraffin (3 g); and (c) polyethylene glycol (5 g), 1,3-butylene glycol (4 g), and distilled water (an amount making total 100 g)

30 Preparation method:

Component(c) was dissolved and adjusted to about 85°C.  
To the solution was added under stirring a material in which component(b) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was  
35 cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 62) Lotion

Prescription:

The lotion is composed of (a) metronidazole (5 g); (b) stearic acid (2 g), cetanol (1.5 g), white petrolatum (4 g), squalane (5 g), caprylic/capric acid triglyceride (2 g), sorbitan monooleate (2 g), polyethylene glycol (5 g); and (c) dipropylene glycol (5 g), triethanol amine (0.7 g), purified water (60 g); (d) isopropanol (10 g), and purified water (an amount making total 100 g)

10 Preparation method:

Component(c) was dissolved and adjusted to about 70°C. To the solution was added under stirring a material in which component(b) had been dissolved and adjusted to about 70°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 40°C while continuously stirring, and then, component(d) was added thereto, and the mixture was cooled to about 25°C under stirring. The resulting lotion was charged in a suitable sealed vessel.

20 (Example 63) Lotion

Prescription:

The lotion is composed of (a) metronidazole (5 g), tranilast (0.4 g); (b) stearic acid (2 g), cetanol (1.5 g), white petrolatum (4 g), squalane (5 g), caprylic/capric acid triglyceride (2 g), sorbitan monooleate (2 g), polyethylene glycol (5 g); and (c) dipropylene glycol (5 g), triethanol amine (0.7 g), purified water (60 g); (d) isopropanol (10 g), and purified water (an amount making total 100 g)

Preparation method:

30 Component(c) was dissolved and adjusted to about 70°C. To the solution was added under stirring a material in which component(b) had been dissolved and adjusted to about 70°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 40°C while continuously stirring, and then, component(d) was added thereto, and the mixture was cooled to about 25°C under stirring. The resulting lotion was charged in a suitable sealed vessel.

(Example 64) Lotion

Prescription:

The lotion is composed of (a) tinidazole (3 g),  
5 clotrimazole (0.1 g), prednisolone acetate (0.005 g); (b)  
stearic acid (2 g), cetanol (1.5 g), white petrolatum (4 g),  
squalane (5 g), caprylic/capric acid triglyceride (2 g),  
sorbitan monooleate (2 g), polyethylene glycol (5 g); and (c)  
dipropylene glycol (5 g), triethanol amine (0.7 g), purified  
10 water (60 g); (d) isopropanol (10 g), and purified water (an  
amount making total 100 g)

Preparation method:

Component(c) was dissolved and adjusted to about 70°C.  
To the solution was added under stirring a material in which  
15 component(b) had been dissolved and adjusted to about 70°C,  
followed by the addition of component(a). The mixture was  
cooled to a temperature of about 40°C while continuously  
stirring, and then, component(d) was added thereto, and the  
mixture was cooled to about 25°C under stirring. The  
20 resulting lotion was charged in a suitable sealed vessel.

(Example 65) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (2 g); (b)  
25 glycol monostearate (10 g), cetanol (7 g), liquid paraffin (9  
g), white petrolatum (3.5 g); and (c) urea (2 g), propylene  
glycol (6.5 g), sodium lauryl sulfate (1 g), and purified  
water (an amount making total 100 g)

Preparation method:

30 Component(b) was dissolved and adjusted to about 85°C.  
To the solution was added under stirring a material in which  
component(c) had been dissolved and adjusted to about 85°C,  
followed by the addition of component(a). The mixture was  
cooled to a temperature of about 25°C while continuously  
35 stirring, and then, charged in a suitable vessel.

(Example 66) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (2 g); (b) glycol monostearate (10 g), cetanol (7 g), liquid paraffin (9 g), white petrolatum (3.5 g); and (c) urea (2 g), propylene glycol (6.5 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

15 (Example 67) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (10 g); (b) glycol monostearate (10 g), cetanol (7 g), liquid paraffin (9 g), white petrolatum (2.5 g); and (c) urea (2 g), polyethylene glycol (7 g), Tween 80 (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

30 (Example 68) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (10 g); (b) glycol monostearate (10 g), cetanol (7 g), liquid paraffin (9 g), white petrolatum (2.5 g); and (c) urea (2 g), polyethylene glycol (7 g), Tween 80 (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 69) Ointment for external use

Prescription:

10 The ointment is composed of (a) metronidazole (3 g);  
(b) white petrolatum (45 g) cetanol (20 g), polyoxyethylene  
hydrogenated castor oil (5 g), liquid paraffin (5 g), propyl  
parahydroxybenzoate (0.1 g); and (c) methyl  
parahydroxybenzoate (0.1 g), Tween 80 (2 g), polyethylene  
15 glycol (5 g), and distilled water (an amount making total 100  
g)

Preparation method:

Comonent(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

25 (Example 70) Ointment for external use

Prescription:

The ointment is composed of (a) tinidazole (3 g); (b) white petrolatum (45 g), cetanol (20 g), polyoxyethylene hydrogenated castor oil (5 g), liquid paraffin (5 g), propyl  
30 parahydroxybenzoate (0.1 g); and (c) methyl  
parahydroxybenzoate (0.1 g), Tween 80 (2 g), polyethylene  
glycol (5 g), and distilled water (an amount making total 100  
g)

Preparation method:

35 Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C,

followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

5 (Example 71) Lotion

Prescription:

The lotion is composed of (a) metronidazole (3 g); (b) stearic acid (2 g), cetanol (1.5 g), white petrolatum (4 g), squalane (5 g), caprylic/caproic acid triglyceride (2 g),  
10 sorbitan monooleate (2 g); (c) polyethylene glycol (5 g), dipropylene glycol (5 g), triethanol amine (0.2 g), purified water (60 g); and (d) isopropanol (10 g), and purified water (an amount making total 100 g)

Preparation method:

15 Component(c) was dissolved and adjusted to about 70°C. To the solution was added under stirring a material in which component(b) had been dissolved and adjusted to about 70°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 40°C while continuously  
20 stirring, and then, component(d) was added thereto, and the mixture was cooled to about 25°C under stirring. The resulting lotion was charged in a suitable sealed vessel.

(Example 72) Lotion

25 Prescription.

The lotion is composed of (a) tinidazole (3 g); (b) stearic acid (2 g), cetanol (1.5 g), white petrolatum (4 g), squalane (5 g), caprylic/caproic acid triglyceride (2 g), sorbitan monooleate (2 g); (c) polyethylene glycol (5 g),  
30 dipropylene glycol (5 g), triethanol amine (0.2 g), purified water (60 g); and (d) isopropanol (10 g), and purified water (an amount making the total amount 100 g)

Preparation method:

Component(c) was dissolved and adjusted to about 70°C.  
35 To the solution was added under stirring a material in which component(b) had been dissolved and adjusted to about 70°C, followed by the addition of component(a). The mixture was

cooled to a temperature of about 40°C while continuously stirring, and then, component(d) was added thereto, and the mixture was cooled to about 25°C under stirring. The resulting lotion was charged in a suitable sealed vessel.

5

(Example 73) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (2 g),  
tranilast (0.1 g); (b) glycol monostearate (5 g),  
10 polyoxyethylene (23) cetyl ether (2 g), stearic acid (0.5 g),  
cetanol (5 g), white petrolatum (3.5 g), liquid paraffin (5  
g), isopropyl myristate (5 g), octyl dodecyl myristate (3 g),  
propyl parahydroxybenzoate (0.15 g); and (c) propylene glycol  
(7 g), methyl parahydroxybenzoate (0.15 g), and distilled  
15 water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 75°C.  
To the solution was added under stirring a material in which  
component(c) had been dissolved and adjusted to about 75°C,  
20 followed by the addition of component(a). The mixture was  
cooled to a temperature of about 25°C while continuously  
stirring, and then, charged in a suitable vessel.

(Example 74) Cream for external use

25 Prescription:

The cream is composed of (a) tinidazole (2 g),  
tranilast (0.1 g); (b) glycol monostearate (5 g),  
polyoxyethylene (23) cetyl ether (2 g), stearic acid (0.5 g),  
cetanol (5 g), white petrolatum (3.5 g), liquid paraffin (5  
30 g), isopropyl myristate (5 g), octyl dodecyl myristate (3 g),  
propyl parahydroxybenzoate (0.15 g); and (c) propylene glycol  
(7 g), methyl parahydroxybenzoate (0.15 g), and distilled  
water (an amount making total 100 g)

Preparation method:

35 Component(b) was dissolved and adjusted to about 75°C.  
To the solution was added under stirring a material in which  
component(c) had been dissolved and adjusted to about 75°C,

followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

5 (Example 75) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (1.5 g); (b) stearic acid (5 g), cetanol (5 g), polyoxyethylene stearyl ether (3 g); and (c) glycerin (6 g), 1,3-butylene glycol (4  
10 g), triethanol amine (0.3 g), and purified water (an amount making total 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added  
15 component(c) heated to about 80°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 75°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a  
20 suitable vessel.

(Example 76) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (3.0 g); (b)  
25 stearic acid (5 g), cetanol (5 g), polyoxyethylene stearyl ether (3 g); and (c) glycerin (6 g), 1,3-butylene glycol (4 g), triethanol amine (0.3 g), and purified water (an amount making total 100 g)

Preparation method:

30 Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 80°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 75°C whereby the mixture was emulsified under  
35 stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a suitable vessel.

(Example 77) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (1.5 g); (b)  
5 stearic acid (5 g), cetanol (5 g), polyoxyethylene stearyl  
ether (3 g); and (c) glycerin (6 g), 1,3-butylene glycol (4  
g), triethanol amine (0.3 g), and purified water (an amount  
making total 100 g)

Preparation method:

10 Component(a) was dissolved in a suitable amount of  
purified water under heating. Then, to the mixture was added  
component(c) heated to about 80°C, and the resulting mixture  
was added to a solution of component(b) dissolved by heating  
at about 75°C whereby the mixture was emulsified under  
15 stirring. The emulsion was cooled to a temperature of about  
30°C while continuously stirring, and then, charged in a  
suitable vessel.

(Example 78) Cream for external use

20 Prescription:

The cream is composed of (a) tinidazole (3.0 g); (b)  
stearic acid (5 g), cetanol (5 g), polyoxyethylene stearyl  
ether (3 g); and (c) glycerin (6 g), 1,3-butylene glycol (4  
g), triethanol amine (0.3 g), and purified water (an amount  
25 making total 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of  
purified water under heating. Then, to the mixture was added  
component(c) heated to about 80°C, and the resulting mixture  
30 was added to a solution of component(b) dissolved by heating  
at about 75°C whereby the mixture was emulsified under  
stirring. The emulsion was cooled to a temperature of about  
30°C while continuously stirring, and then, charged in a  
suitable vessel.

35 (Example 79) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (1.0 g); (b) stearic acid (0.5 g), glycol monostearate (8 g), stearyl alcohol (5 g), liquid paraffin (8 g); and (c) propylene glycol (6 g), glycerin (4 g), sodium polyoxyethylene lauryl ether sulfate (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component (a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component (c) heated to about 80°C, and the resulting mixture was added to a solution of component (b) dissolved by heating at about 75°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a suitable vessel.

(Example 80) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (2.5 g); (b) stearic acid (0.5 g), glycol monostearate (8 g), stearyl alcohol (5 g), liquid paraffin (8 g); and (c) propylene glycol (6 g), glycerin (4 g), sodium polyoxyethylene lauryl ether sulfate (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component (a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component (c) heated to about 80°C, and the resulting mixture was added to a solution of component (b) dissolved by heating at about 75°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a suitable vessel.

(Example 81) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (1.0 g); (b)

stearic acid (0.5 g), glycol monostearate (8 g), stearyl alcohol (5 g), liquid paraffin (8 g); and (c) propylene glycol (6 g), glycerin (4 g), sodium polyoxyethylene lauryl ether sulfate (1 g), and purified water (an amount making  
5 total 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 80°C, and the resulting mixture  
10 was added to a solution of component(b) dissolved by heating at about 75°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a suitable vessel.

15 (Example 82) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (2.5 g); (b) stearic acid (0.5 g), glycol monostearate (8 g), stearyl  
20 alcohol (5 g), liquid paraffin (8 g); and (c) propylene glycol (6 g), glycerin (4 g), sodium polyoxyethylene lauryl ether sulfate (1 g), and purified water (an amount making total 100 g)

Preparation method:

25 Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component (c) heated to about 80°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 75°C whereby the mixture was emulsified under  
30 stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a suitable vessel.

(Example 83) Cream for external use

35 Prescription:

The cream is composed of (a) metronidazole (1.0 g); (b) glycol monostearate (10.4 g), cetanol (7.3 g), liquid

paraffin (9 g), white petrolatum (3.5 g), propylparaben (0.05 g); and (c) propylene glycol (6.5 g), methylparaben (0.05 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

5 Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 80°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 75°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a suitable vessel.

15 (Example 84) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (1.8 g); (b) glycol monostearate (10.4 g), cetanol (7.3 g), liquid paraffin (9 g), white petrolatum (3.5 g), propylparaben (0.05 g); and (c) propylene glycol (6.5 g), methylparaben (0.05 g), sodium lauryl sulfate (1 g), the purified water (an amount making total 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 80°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 75°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a suitable vessel.

(Example 85) Cream for external use

Prescription:

35 The cream is composed of (a) tinidazole (1.0 g); (b) glycol monostearate (10.4 g), cetanol (7.3 g), liquid paraffin (9 g), white petrolatum (3.5 g), propylparaben (0.05

g); and (c) propylene glycol (6.5 g), methylparaben (0.05 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

Preparation method:

5       Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 80°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 75°C whereby the mixture was emulsified under  
10   stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a suitable vessel.

(Example 86) Cream for external use

15   Prescription:

      The cream is composed of (a) tinidazole (2.0 g); (b) glycol monostearate (10.4 g), cetanol (7.3 g), liquid paraffin (9 g), white petrolatum (3.5 g), propylparaben (0.05 g); and (c) propylene glycol (6.5 g), methylparaben (0.05 g),  
20   sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

Preparation method:

      Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added  
25   component(c) heated to about 80°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 75°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a  
30   suitable vessel.

(Example 87) Cream for external use

Prescription:

      The cream is composed of (a) tinidazole (1.0 g); (b)  
35   glycol monostearate (10.4 g), cetanol (7.3 g), liquid paraffin (9 g), white petrolatum (3.5 g), propylparaben (0.05 g); and (c) propylene glycol (6.5 g), methylparaben (0.05 g),

sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 85°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 85°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 88) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (2.0 g); (b) glycol monostearate (10.4 g), cetanol (7.3 g), liquid paraffin (9 g), white petrolatum (3.5 g), propylparaben (0.05 g); and (c) propylene glycol (6.5 g), methylparaben (0.05 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 85°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 85°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 89) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (1.8 g); (b) glycol monostearate (10.4 g), cetanol (7.3 g), liquid paraffin (9 g), white petrolatum (3.5 g), propylparaben (0.05 g); and (c) propylene glycol (6.5 g), methylparaben (0.05 g), sodium lauryl sulfate (1 g), and purified water (an amount

making total 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added  
5 component(c) heated to about 85°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 85°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a  
10 suitable vessel.

(Example 90) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (1.5 g); (b)  
15 glycol monostearate (10.4 g), cetanol (7.3 g), liquid paraffin (9 g), white petrolatum (3.5 g), propylparaben (0.05 g); and (c) propylene glycol (6.5 g), methylparaben (0.05 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

20 Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 85°C, and the resulting mixture was added to a solution of component(b) dissolved by heating  
25 at about 85°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

30 (Example 91) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (2.0 g); (b)  
glycol monostearate (7 g), stearyl alcohol (7 g), liquid paraffin (5 g), polyoxyethylene cetostearyl ether (3 g); and  
35 (c) glycerin (5 g), 1,3-butylene glycol (7 g), sodium carboxymethyl cellulose (0.4 g), Tween 80 (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 85°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 80°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 92) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (2.0 g); (b) glycol monostearate (7 g), stearyl alcohol (7 g), liquid paraffin (5 g), polyoxyethylene cetostearyl ether (3 g); and (c) glycerin (5 g), 1,3-butylene glycol (7 g), sodium carboxymethyl cellulose (0.4 g), Tween 80 (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 85°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 80°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 93) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (5.0 g); (b) glycol monostearate (7 g), stearyl alcohol (4 g), white petrolatum (3.5 g), isopropyl myristate (3 g), Span 60 (1 g), Tween 60 (0.5 g); and (c) propylene glycol (7 g), glycerin (2 g), Tween 80 (0.1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 80°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 75°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a suitable vessel.

10 (Example 94) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (3.0 g); (b) glycol monostearate (7 g), stearyl alcohol (4 g), white petrolatum (3.5 g), isopropyl myristate (3 g), Span 60 (1 g), Tween 60 (0.5 g); and (c) propylene glycol (7 g), glycerin (2 g), Tween 80 (0.1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 80°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 75°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a suitable vessel.

(Example 95) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (1.5 g); (b) glycol monostearate (7.28 g), sorbitan monostearate (3.12 g), cetanol (7.3 g), white petrolatum (3.5 g), liquid paraffin (9 g), propylparaben (0.05 g); and (c) propylene glycol (6.5 g), sodium lauryl sulfate (1 g), methylparaben (0.05 g), and purified water (an amount making the total amount 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of

purified water under heating. Then, to the mixture was added component(c) heated to about 80°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 85°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a suitable vessel.

(Example 96) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (2.0 g); (b) glycol monostearate (7.28 g), sorbitan monostearate (3.12 g), cetanol (7.3 g), white petrolatum (3.5 g), liquid paraffin (9 g), propylparaben (0.05 g); and (c) propylene glycol (6.5 g), sodium lauryl sulfate (1 g), methylparaben (0.05 g), and purified water (an amount making total 100 g)

Preparation method:

Component(a) was dissolved in a suitable amount of purified water under heating. Then, to the mixture was added component(c) heated to about 80°C, and the resulting mixture was added to a solution of component(b) dissolved by heating at about 85°C whereby the mixture was emulsified under stirring. The emulsion was cooled to a temperature of about 30°C while continuously stirring, and then, charged in a suitable vessel.

(Example 97) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (2.0 g); (b) glycerol monostearate (6 g), stearyl alcohol (5 g), cetanol (6 g), isopropyl myristate (1 g), Span 60 (1.5 g), Tween 60 (1 g); and (c) sodium carboxymethyl cellulose (0.2 g), propylene glycol (4 g), and purified water (an amount making the total amount 100 g)

Preparation method:

Component(b) was dissolved under heating and adjusted to about 75°C. To the solution was added under stirring a

material in which component(c) had been dissolved and adjusted to about 75°C, followed by the addition of component(a) and stirring. The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 98) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (1.0 g); (b) glycol monostearate (7 g), stearyl alcohol (4 g), white petrolatum (3 g), isopropyl myristate (3 g), Span 60 (1 g), Tween 60 (0.5 g); and (c) propylene glycol (7 g), glycerin (2 g), Tween 80 (0.1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved under heating and adjusted to about 75°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 75°C, followed by the addition of component(a) and stirring. The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 99) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (1.0 g); (b) glycol monostearate (7 g), stearyl alcohol (4 g), white petrolatum (3 g), isopropyl myristate (3 g), Span 60 (1 g), Tween 60 (0.5 g); and (c) propylene glycol (7 g), glycerin (2 g), Tween 80 (0.1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved under heating and adjusted to about 75°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 75°C, followed by the addition of component(a) and stirring. The mixture was cooled to a

temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 100) Cream for external use

5 Prescription:

The cream is composed of (a) metronidazole (2.0 g),  
nofloxacin (0.05 g); (b) glyceryl monostearate (2 g), stearyl  
alcohol (5 g), white petrolatum (3 g), isopropyl myristate (3  
10 (7 g), glycerin (2 g), Tween 60 (0.5 g); and (c) propylene glycol  
(an amount making total 100 g)

Preparation method:

Component(b) was dissolved under heating and adjusted  
to about 75°C. To the solution was added under stirring a  
15 material in which component(c) had been dissolved and  
adjusted to about 75°C, followed by the addition of  
component(a) and stirring. The mixture was cooled to a  
temperature of about 25°C while continuously stirring, and  
then, charged in a suitable vessel.

20

(Example 101) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (2.0 g),  
tranilast (0.2 g); (b) glycol monostearate (4 g), cetanol (4  
25 g), stearyl alcohol (3 g), polyoxyethylene cetyl alcohol (2  
g), isopropyl myristate (3 g), Span 60 (1 g), Tween 60 (0.5  
g); and (c) propylene glycol (5 g), and purified water (an  
amount making total 100 g)

Preparation method:

30 Component(b) was dissolved under heating and adjusted  
to about 75°C. To the solution was added under stirring a  
material in which component(c) had been dissolved and  
adjusted to about 75°C, followed by the addition of  
component(a) and stirring. The mixture was cooled to a  
35 temperature of about 25°C while continuously stirring, and  
then, charged in a suitable vessel.

(Example 102) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (2.0 g),  
ketoprofen (0.5 g); (b) glycol monostearate (4 g), cetanol (4  
5 g), stearyl alcohol (3 g), polyoxyethylene cetyl alcohol (2  
g), isopropyl myristate (3 g), Span 60 (1 g), Tween 60 (0.5  
g); and (c) propylene glycol (5 g), and purified water (an  
amount making total 100 g)

Preparation method:

10 Component(b) was dissolved under heating and adjusted  
to about 75°C. To the solution was added under stirring a  
material in which Component(c) had been dissolved and  
adjusted to about 75°C, followed by the addition of  
component(a) and stirring. The mixture was cooled to a  
15 temperature of about 25°C while continuously stirring, and  
then, charged in a suitable vessel.

(Example 103) Cream for external use

Prescription:

20 The cream is composed (a) metronidazole (2.5 g),  
procaine hydrochloride (0.2 g); (b) glycerol monostearate (2  
g), stearyl alcohol (5 g), white petrolatum (3 g), isopropyl  
myristate (3 g), Span 60 (1 g), Tween 60 (0.5 g); and (c)  
propylene glycol (7 g), glycerin (2 g), Tween 80 (0.1 g), and  
25 purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved under heating and adjusted  
to about 75°C. To the solution was added under stirring a  
material in which component(c) had been dissolved and  
30 adjusted to about 75°C, followed by the addition of  
component(a) and stirring. The mixture was cooled to a  
temperature of about 25°C while continuously stirring, and  
then, charged in a suitable vessel.

35 (Example 104) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (3.0 g),

tamoxifen citrate (0.05 g); (b) glycol monostearate (4 g),  
cetanol (4 g), stearyl alcohol (3 g), polyoxyethylene cetyl  
alcohol (2 g), isopropyl myristate (3 g), Span 60 (1 g),  
Tween 60 (0.5 g); and (c) propylene glycol (5 g), and  
5 purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved under heating and adjusted  
to about 75°C. To the solution was added under stirring a  
material in which component(c) had been dissolved and  
10 adjusted to about 75°C, followed by the addition of  
component(a) and stirring. The mixture was cooled to a  
temperature of about 25°C while continuously stirring, and  
then, charged in a suitable vessel.

15 (Example 105) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (2.0 g),  
carpronium chloride (0.5 g); (b) stearic acid (0.5 g), glycol  
monostearate (12 g), stearyl alcohol (7 g), white petrolatum  
20 (2 g), liquid paraffin (5 g); and (c) polyethylene glycol (5  
g), 1,3-butylene glycol (5 g), and purified water (an amount  
making total 100 g)

Preparation method:

Component(b) was dissolved under heating and adjusted  
25 to about 75°C. To the solution was added under stirring a  
material in which component(c) had been dissolved and  
adjusted to about 75°C, followed by the addition of  
component(a) and stirring. The mixture was cooled to a  
temperature of about 25°C while continuously stirring, and  
30 then, charged in a suitable vessel.

(Example 106) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (2.0 g),  
35 extract of calves blood (0.5 g); (b) stearic acid (0.5 g),  
glycol monostearate (12 g), stearyl alcohol (7 g), white  
petrolatum (2 g), liquid paraffin (5 g); and (c) polyethylene

glycol (5 g), 1,3-butylene glycol (5 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved under heating and adjusted to about 75°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 75°C, followed by the addition of component(a) and stirring. The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 107) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (0.5 g); (b) stearic acid (0.5 g), glycol monostearate (10 g), cetanol (5 g), white petrolatum (3 g); and (c) propylene glycol (7 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved under heating and adjusted to about 75°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 75°C, followed by the addition of component(a) and stirring. The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 108) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (0.5 g); (b) stearic acid (0.5 g), glycol monostearate (10 g), cetanol (5 g), white petrolatum (3 g); and (c) propylene glycol (7 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved under heating and adjusted to about 75°C. To the solution was added under stirring a

material in which Component(c) had been dissolved and adjusted to about 75°C, followed by the addition of component(a) and stirring. The mixture was cooled to a temperature of about 25°C while continuously stirring, and  
5 then, charged in a suitable vessel.

(Example 109) Cream for external use

Prescription:

10 The cream is composed of (a) metronidazole (5 g); (b) stearic acid (0.5 g), glycol monostearate (10 g), cetanol (5 g), white petrolatum (3 g); and (c) propylene glycol (7 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

Preparation method:

15 Component(b) was dissolved under heating and adjusted to about 75°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 75°C, followed by the addition of component(a) and stirring. The mixture was cooled to a  
20 temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 110) Cream for external use

Prescription:

25 The cream is composed of (a) tinidazole (5 g); (b) stearic acid (0.5 g), glycol monostearate (10 g), cetanol (5 g), white petrolatum (3 g); and (c) propylene glycol (7 g), sodium lauryl sulfate (1 g), and purified water (an amount making total 100 g)

30 Preparation method:

Component(b) was dissolved under heating and adjusted to about 75°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 75°C, followed by the addition of  
35 component(a) and stirring. The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 111) Ointment

Prescription:

The ointment is composed of (a) metronidazole (2 g);  
5 (b) stearic acid (2 g), glycol monostearate (12 g),  
polyoxyethylene glycol monostearate (3 g), polyoxyethylene  
cetyl/stearyl ether (12E.O.) (1 g), polyoxyethylene  
cetyl/stearyl ether (20E.O.) (1 g), cetanol (2 g), liquid  
paraffin (8 g); and (c) 1,3-butylene glycol (7 g), glycerin  
10 (5 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 75°C.  
To the solution was added under stirring a material in which  
Component(c) had been dissolved and adjusted to about 75°C,  
15 followed by the addition of component(a). The mixture was  
cooled to a temperature of about 25°C while continuously  
stirring, and then, charged in a suitable vessel.

(Example 112) External preparation

20 Prescription:

The preparation is composed of (a) metronidazole (2 g),  
ketoconazole (0.2 g); (b) glyceryl monostearate (7.5 g),  
sorbitan monostearate (3 g), stearyl alcohol (7 g), liquid  
paraffin (8 g), white petrolatum (5 g), span 80 (1 g); and  
25 (c) propylene glycol (5 g), 1,3-butylene glycol (3 g), Tween  
80 (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 75°C.  
To the solution was added under stirring a material in which  
30 component(c) had been dissolved and adjusted to about 75°C,  
followed by the addition of component(a). The mixture was  
cooled to a temperature of about 25°C while continuously  
stirring, and then, charged in a suitable vessel.

35 (Example 113) External preparation

Prescription:

The preparation is composed of (a) metronidazole (2 g);

(b) glyceryl monostearate (7.5 g), sorbitan monostearate (3 g), stearyl alcohol (7 g), liquid paraffin (8 g), white petrolatum (5 g), span 80 (1 g); and (c) propylene glycol (5 g), 1,3-butylene glycol (3 g), Tween 80 (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 75°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 75°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 114) External preparation

Prescription:

The preparation is composed of (a) tinidazole (2 g), isoconazole nitrate (0.2 g); (b) glycol monostearate (7.5 g), sorbitan monostearate (3 g), stearyl alcohol (7 g), liquid paraffin (8 g), white petrolatum (5 g), span 80 (1 g); and (c) propylene glycol (5 g), 1,3-butylene glycol (3 g), Tween 80 (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 75°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 75°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 115) External preparation

Prescription:

The preparation is composed of (a) tinidazole (2 g); (b) glyceryl monostearate (7.5 g), sorbitan monostearate (3 g), stearyl alcohol (7 g), liquid paraffin (8 g), white petrolatum (5 g), span 80 (1 g); and (c) propylene glycol (5 g), 1,3-butylene glycol (3 g), Tween 80 (1 g), purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 75°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 75°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 116) Shampoo

Prescription:

The shampoo is composed of tinidazole (1.5 g), polyglyceryl monolaurate (4 g), sodium polyoxyethylene lauryl ether sulfate (7 g), lauryldimethylaminoacetate betaine (2.5 g), coconut fatty acid diethanol amide (4 g), polyethylene glycol (5 g), 1,3-butylene glycol (3 g), citric acid (a suitable amount), and purified water (an amount making total 100 g)

Preparation method:

Metronidazole was added to a mixture of suitable amounts of polyethylene glycol and purified water, and the mixture was melted by heating. In other vessel were weighed suitable amounts of polyglyceryl monolaurate, sodium polyoxyethylene lauryl ether sulfate, lauryl dimethylacetate betaine, coconut fatty acid diethanol amide, polyethylene glycol, 1,3-butylene glycol and purified water, and the mixture was heated to about 70°C under stirring and added to a mixture of tinidazole, polyethylene glycol and purified water. A pH of the mixture was adjusted to about 6.5 with citric acid. The resulting mixture was cooled until a temperature thereof became about 25°C under stirring.

(Example 117) Rinse

Prescription:

The rinse is composed of (a) metronidazole (2 g); (b) isopropyl myristate (1 g), butyl myristate (1 g), silicone oil (2 g), liquid paraffin (1 g), a hydrochloric acid solution of N-[alkyl(12,14)oxy-2-hydroxypropyl]-L-alginic

acid (2 g); and (c) lactic acid (0.05 g), polyethylene glycol (6 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was heated to about 80°C, and to the mixture was added a material in which component(a) had been added to component(c) while stirring to melt the material under heating and adjusted to about 80°C. The mixture was cooled to about 25°C under stirring and charged in a suitable vessel.

(Example 118) Rinse

Prescription:

The rinse is composed of (a) tinidazole (2 g); (b) isopropyl myristate (1 g), butyl myristate (1 g), silicone oil (2 g), liquid paraffin (1 g), a hydrochloric acid solution of N-[alkyl(12,14)oxy-2-hydroxypropyl]-L-alginic acid (2 g); and (c) lactic acid (0.05 g), polyethylene glycol (6 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was heated to about 80°C, and to the mixture was added a material in which component(a) had been added to component(c) while stirring to melt the material under heating and adjusted to about 80°C. The mixture was cooled to about 25°C under stirring and charged in a suitable vessel.

(Example 119) Soap

Prescription:

The soap is composed of metronidazole (3 g), monoglycerol laurate (75 g), sodium monoglyceryl fatty acid sulfate (7 g), stearyl alcohol (8 g), silicone oil (1 g), glycerin (3 g), polyethylene glycol (5 g), sodium carboxymethyl cellulose (0.4 g), purified water (an amount making total 100 g), and perfume (a suitable amount)

Preparation method:

Compositions except perfume were melted by heating under stirring. Cooling was initiated and perfume was added

before the melt was solidified. The solid material was dried in a dark place with a sufficient time to obtain soap.

(Example 120) Soap

5 Prescription:

The soap is composed of (a) tinidazole (2 g); (b) monoglycerol laurate (75 g), sodium monoglyceryl fatty acid sulfate (7 g), stearyl alcohol (8 g), silicone oil (1 g), glycerin (3 g), polyethylene glycol (5 g), sodium  
10 carboxymethyl cellulose (0.4 g), purified water (an amount making total 100 g), and perfume (a suitable amount)

Preparation method:

Compositions except perfume were melted by heating under stirring. Cooling was initiated and perfume was added  
15 before the melt was solidified. The solid material was dried in a dark place with a sufficient time to obtain soap.

(Example 121) Face lotion

Prescription:

20 The lotion is composed of (a) metronidazole (1 g); (b) propylene glycol (3 g), polyethylene glycol (5 g), sodium carboxymethyl cellulose (0.4 g); (c) polyoxethylen oleyl cetyl ether (1 g), jojoba oil (0.5 g); (d) perfume (a suitable amount), ethanol (8 g); and (e) purified water (an amount  
25 making total 100 g)

Preparation method:

Component(b) was added to (e) and the mixture was melted by heating. To the above mixture was added component(a) and the resulting mixture was melted and cooled  
30 to room temperature. Further, to the above mixture was added a material in which component(c) had been dissolved and dispersed in component(d) and the resulting mixture was stirred and homogenized.

35 (Example 122) Face lotion

Prescription:

The lotion is composed of (a) tinidazole (0.5 g); (b)

propylene glycol (3 g), polyethylene glycol (5 g), sodium carboxymethyl cellulose (0.4 g); (c) polyoxethylene oleyl cetyl ether (1 g), jojoba oil (0.5 g); (d) perfume (a suitable amount), ethanol (8 g); and (e) purified water (an amount making total 100 g)

Preparation method:

Component(b) was added to component(e) and the mixture was melted by heating. To the above mixture was added component(a) and the resulting mixture was melted and cooled to room temperature. Further, to the above mixture was added a material in which component(c) had been dissolved and dispersed in component(d) and the resulting mixture was stirred and homogenized.

(Example 123) Gel

Prescription:

The gel is composed of tinidazole (1 g), polyethylene glycol (8 g), carboxyvinyl polymer (0.5 g), methyl cellulose (0.2 g), propylene glycol (5 g), glycerin (2 g), polyoxyethylene oleyl cetyl ether (1 g), isopropanol (5 g), sodium hydroxide (a suitable amount), an purified water (an amount making total 100 g)

Preparation method:

Polyethylene glycol was added to purified water and melted, and after tinidazole was added to the mixture, the mixture was dissolved under heating. The solution was cooled to about 50°C, and to the solution were added under stirring a material in which polyoxyethylene cetyl ether had been added to propylene glycol and glycerin heated to about 50°C. Further, under continuous stirring, sodium hydroxide was added to the above mixture and a pH thereof was adjusted to about 6.8. After cooling the resulting mixture to about 40°C, isopropanol was added thereto, and the resulting mixture was cooled to about 25°C, and charged in a suitable vessel.

(Example 124) Cream

Prescription:

The cream is composed of (a) secnidazole (2 g); (b) glycol monostearate (10 g), polyoxyethylene glycol monostearate (3 g), polyoxyethylene cetyl/stearyl ether (2 g), cetanol (4 g), beeswax (1 g), octyl dodecyl myristate (7 g), isopropyl myristate (2 g); (c) polyethylene glycol (3 g), carboxyvinyl polymer (0.2 g), purified water (an amount making total 100 g); and (d) an aqueous sodium hydroxide solution (a suitable amount)

Preparation method:

Component(b) was heated to about 75°C, then, to component(b) was added component(c) heated to about 75°C under stirring, followed by the addition of component(a) under stirring. Thereafter, a pH of the mixture was adjusted to about 6.8 with component(d). Thereafter, the resulting mixture was cooled to a temperature of about 25°C, and the resulting cream was charged in a suitable vessel.

(Example 125) Cream

Prescription:

The cream is composed of (a) Panidazole (2 g); (b) glycol monostearate (10 g), polyoxyethylene glycol monostearate (3 g), polyoxyethylene cetyl/stearyl ether (2 g), cetanol (4 g), beeswax (1 g), octyldodecyl myristate (7 g), isopropyl myristate (2 g); (c) polyethylene glycol (3 g), carboxyvinyl polymer (0.2 g), purified water (an amount making total 100 g); and (d) an aqueous sodium hydroxide solution (a suitable amount)

Preparation method:

Component(b) was heated to about 75°C, then, to component(b) was added component(c) heated to about 75°C under stirring, followed by the addition of component(a) under stirring. Thereafter, a pH of the mixture was adjusted to about 6.8 with (d). Thereafter, the resulting mixture was cooled to a temperature of about 25°C, and the resulting cream was charged in a suitable vessel.

(Example 126) Cream

Prescription:

The cream is composed of (a) dimetridazole (2 g); (b) glycol monostearate (10 g), polyoxyethylene glycol monostearate (3 g), polyoxyethylene cetyl/stearyl ether (2 g),  
5 cetanol (4 g), beeswax (1 g), octyl dodecyl myristate (7 g), isopropyl myristate (2 g); (c) polyethylene glycol (3 g), carboxyvinyl polymer (0.2 g), purified water (an amount making total 100 g); (d) an aqueous sodium hydroxide solution (a suitable amount)

10 Preparation method:

Component(b) was heated to about 75°C, then, to component(b) was added component(c) heated to about 75°C under stirring, followed by the addition of component(a) under stirring. Thereafter, a pH of the mixture was adjusted  
15 to about 6.8 with component(d). Thereafter, the resulting mixture was cooled to a temperature of about 25°C, and the resulting cream was charged in a suitable vessel.

(Example 127) Cream

20 Prescription:

The cream is composed of (a) ronidazole(2 g); (b) glycol monostearate (10 g), polyoxyethylene glycol monostearate (3 g), polyoxyethylene cetostearyl ether (2 g),  
25 cetanol (4 g), beeswax (1 g), octyl dodecyl myristate (7 g), isopropyl myristate (2 g); (c) polyethylene glycol (3 g), carboxyvinyl polymer (0.2 g), purified water (an amount making total 100 g); and (d) an aqueous sodium hydroxide solution (a suitable amount)

Preparation method:

30 (b) was heated to about 75°C, then, to (b) was added (c) heated to about 75°C under stirring, followed by the addition of (a) under stirring. Thereafter, a pH of the mixture was adjusted to about 6.8 with (d). Thereafter, the resulting mixture was cooled to a temperature of about 25°C,  
35 and the resulting cream was charged in a suitable vessel.

(Example 128) Cream

Prescription:

The cream is composed of (a) ipronidazole (2 g); (b) glycol monostearate (10 g), polyoxyethylene glycol monostearate (3 g), polyoxyethylene cetyl/stearyl ether (2 g),  
5 cetanol (4 g), beeswax (1 g), octyl dodecyl myristate (7 g), isopropyl myristate (2 g); (c) polyethylene glycol (3 g), carboxyvinyl polymer (0.2 g), purified water (an amount making the total amount 100 g); (d) an aqueous sodium hydroxide solution (a suitable amount)

10 Preparation method:

Component(b) was heated to about 75°C, then, to component(b) was added component(c) heated to about 75°C under stirring, followed by the addition of (a) under stirring. Thereafter, a pH of the mixture was adjusted to  
15 about 6.8 with component(d). Thereafter, the resulting mixture was cooled to a temperature of about 25°C, and the resulting cream was charged in a suitable vessel.

(Example 129) Cream

20 Prescription:

The cream is composed of (a) ornidazole (2 g); (b) glycol monostearate (10 g), polyoxyethylene glycol monostearate (3 g), polyoxyethylene cetyl/stearyl ether (2 g),  
25 cetanol (4 g), beeswax (1 g), octyl dodecyl myristate (7 g), isopropyl myristate (2 g); and (c) polyethylene glycol (3 g), carboxyvinyl polymer (0.2 g), purified water (an amount making total 100 g); and (d) an aqueous sodium hydroxide solution (a suitable amount)

Preparation method:

30 Component(b) was heated to about 75°C, then, to component(b) was added component(c) heated to about 75°C under stirring, followed by the addition of component(a) under stirring. Thereafter, a pH of the mixture was adjusted to about 6.8 with component(d). Thereafter, the resulting  
35 mixture was cooled to a temperature of about 25°C, and the resulting cream was charged in a suitable vessel.

(Example 130) Cream for external use

Prescription:

The cream is composed of (a) metronidazole (5 g); (b) glycol monostearate (10 g), cetanol (7 g), liquid praffin (9 g), white petrolatum (2.5 g); and (c) urea (2 g), polyethylene glycol (7 g), Tween 80 (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

(Example 131) Cream for external use

Prescription:

The cream is composed of (a) tinidazole (5 g); (b) glycol monostearate (10 g), cetanol (7 g), liquid praffin (9 g), white petrolatum (2.5 g); and (c) urea (2 g), polyethylene glycol (7 g), Tween 80 (1 g), and purified water (an amount making total 100 g)

Preparation method:

Component(b) was dissolved and adjusted to about 85°C. To the solution was added under stirring a material in which component(c) had been dissolved and adjusted to about 85°C, followed by the addition of component(a). The mixture was cooled to a temperature of about 25°C while continuously stirring, and then, charged in a suitable vessel.

[Test Examples]

(Test Example 1) Treatment of Atopic Dermatitis

The therapeutic effect of the ointment produced in the above Example 1 was examined by applying to patients with atopic dermatitis.

The ointment was applied to the target patients indicated below.

Target patient A: 1-year-old male infant suffering from atopic dermatitis

Target patient B: 2-year-old male infant suffering from atopic dermatitis

5 Target patient C: 40-year-old female suffering from atopic dermatitis

Target patient D: 60-year-old female suffering from atopic dermatitis

10 Target patient E: 27-year-old male suffering from atopic dermatitis

For target patients A and B, the ointment for external use produced in Example 1 was applied twice a day for 4 consecutive weeks to the face appearing prominent atopic dermatitis, and the status of inflammation was observed.

15 In addition, for target patients C, D and E, the ointment for external use produced in the Example 1 was applied twice a day for 4 consecutive weeks to affected areas of prominent atopic dermatitis extending from the lower leg to the ankle, and the status of inflammation was observed.

20 Therapeutic effects were evaluated by scoring rash, eczema and other dermatitis symptoms at the start of treatment along with the status of healing over 3 days, 1 week, 2 weeks, 3 weeks and 4 weeks later. In addition, the presence of itchiness of the skin surface and skin condition  
25 were evaluated after 4 weeks.

Furthermore, evaluation scores were determined in the manner indicated below.

5: Prominent rash, eczema and other dermatitis symptoms, extreme itchiness, subconscious scratching of the skin  
30 surface and the presence of resulting scratches.

4: Prominent rash, eczema and other dermatitis symptoms, itchiness but not to the extend of grade 5.

3: Rash, eczema and other dermatitis symptoms able to be confirmed and only bothersome itchiness.

35 2: Rash, eczema and other dermatitis symptoms only able to be confirmed slightly, and not that much different from normal skin.

1: Absence of rash, eczema and other dermatitis symptoms, no itchiness and appearance of normal skin.

Those results are shown in Table 2 below.

[Table 2]

Patient	Start of treat- ment	After 3 days	After 1 week	After 2 weeks	After 3 weeks	After 4 weeks	Overall evaluation	
							Itchi- ness	Skin surfac e
A	5	5	3	1	1	1	none	Normal
B	5	4	3	2	1	1	none	Normal
C	5	5	4	1	1	1	none	Normal
D	5	5	4	3	2	1	none	Normal
E	3	3	1	1	1	1	none	Normal

As is clear from the above results, the external preparation of the present invention was observed to demonstrate improvement of dermatitis symptoms in 3-7 days after the start of application during treatment of atopic dermatitis, and the skin was no different from completely normal skin 3-4 weeks after the start of application. Furthermore, there was no irritation of the skin by the preparation during application. In addition, there were also no adverse side effects such as rebound observed, which are observed with steroid-based external preparations, even after administration was discontinued.

#### (Test Example 2) Treatment of Atopic Dermatitis

The therapeutic effect of the cream produced in the above Example 4 was examined by applying to patients with atopic dermatitis. The cream was applied to the target patients indicated below.

Target patient F: 2-year-old male infant suffering from atopic dermatitis

Target patient G: 8-year-old male infant suffering from atopic dermatitis

Target patient H: 50-year-old female suffering from atopic dermatitis

Target patient I: 40-year-old female suffering from atopic dermatitis

5 Target patient J: 27-year-old male suffering from atopic dermatitis

For target patients F, G, H, I and J, the cream for external use produced in the Example 4 was applied twice a day for 4 consecutive weeks to the face that exhibited prominent atopic dermatitis, and the status of inflammation was observed.

Therapeutic effects were evaluated by scoring rash, eczema and other dermatitis symptoms at the start of treatment along with the status of healing over 3 days, 1 week, 2 weeks, 3 weeks and 4 weeks later in the same manner as the Test Example 1. In addition, the presence of itchiness of the skin surface and skin condition were evaluated after 4 weeks. Furthermore, the evaluation scores in the above Test Example 1 were used for evaluation.

Those results are shown in Table 3 below.

[Table 3]

Patient	Start of treatment	After 3 days	After 1 week	After 2 weeks	After 3 weeks	After 4 weeks	Overall evaluation	
							Itchiness	Skin surface
F	5	4	2	1	1	1	none	Normal
G	5	4	3	2	1	1	none	Normal
H	5	5	3	1	1	1	none	Normal
I	3	2	2	1	1	1	none	Normal
J	3	2	1	1	1	1	none	Normal

As is clear from the above results, the external preparation of the present invention was observed to demonstrate improvement of dermatitis symptoms in 3-7 days after the start of application during treatment of atopic

dermatitis, and the skin was no different from complete normal skin 3-4 weeks after the start of application. Furthermore, there was no irritation of the skin by the preparation during application. In addition, there were also  
5 no adverse side effects such as rebound observed, which are observed with steroid-based external preparations, even after administration was discontinued.

(Test Example 3) Treatment of Atopic Dermatitis

10 The therapeutic effect of the ointment produced in the above Example 11 was examined by applying to patients with atopic dermatitis. The cream was applied to the target patients indicated below.

15 Target patient K: 1-year-old male infant suffering from atopic dermatitis

Target patient L: 2-year-old male infant suffering from atopic dermatitis

Target patient M: 35-year-old female suffering from atopic dermatitis

20 Target patient N: 54-year-old female suffering from atopic dermatitis

Target patient O: 27-year-old male suffering from atopic dermatitis

25 For target patients K and L, the ointment for external use produced in the Example 11 was applied twice a day for 4 consecutive weeks to the face that exhibited prominent atopic dermatitis, and the status of inflammation was observed.

30 In addition, for target patients M, N and O, the ointment for external use produced in the Example 11 was applied twice a day for 4 consecutive weeks to affected areas of prominent atopic dermatitis extending from the lower leg to the ankle, and the status of inflammation was observed.

35 Therapeutic effects were evaluated by scoring rash, eczema and other dermatitis symptoms at the start of treatment along with the status of healing over 3 days, 1 week, 2 weeks, 3 weeks and 4 weeks later in the same manner as the above Test Example 1. In addition, the presence of

itchiness of the skin surface and skin condition were evaluated after 4 weeks. Furthermore, the evaluation scores in the above Test Example 1 were used for evaluation.

Those results are shown in Table 4 below.

5 [Table 4]

Patient	Start of treatment	After 3 days	After 1 week	After 2 weeks	After 3 weeks	After 4 weeks	Overall evaluation	
							Itchi- ness	Skin surface
K	5	5	3	1	1	1	none	Normal
L	5	4	3	2	1	1	none	Normal
M	5	5	4	1	1	1	none	Normal
N	5	5	4	3	2	1	none	Normal
O	4	3	1	1	1	1	none	Normal

As is clear from the above results, the external preparation of the present invention was observed to demonstrate improvement of dermatitis symptoms in 3-7 days after the start of application during treatment of atopic dermatitis, and the skin was no different from complete normal skin 3-4 weeks after the start of application. Furthermore, there was no irritation of the skin by the preparation during application. In addition, there were also no adverse side effects such as rebound observed, which are observed with steroid-based external preparations, even after administration was discontinued.

#### (Test Example 4) Treatment of Atopic Dermatitis

The therapeutic effect of the cream produced in the above Example 14 was examined by applying to patients with atopic dermatitis.

The cream was applied to the target patients indicated below.

Target patient P: 2-year-old male infant suffering from atopic dermatitis

Target patient Q: 6-year-old male infant suffering from atopic dermatitis

Target patient R: 53-year-old female suffering from atopic dermatitis

5 Target patient S: 58-year-old female suffering from atopic dermatitis

Target patient T: 35-year-old male suffering from atopic dermatitis

10 For target patients P, Q, R, S and T, the cream for external use produced in the Example 14 was applied twice a day for 4 consecutive weeks to the face that exhibited prominent atopic dermatitis, and the status of inflammation was observed.

15 Therapeutic effects were evaluated by scoring rash, eczema and other dermatitis symptoms at the start of treatment along with the status of healing over 3 days, 1 week, 2 weeks, 3 weeks and 4 weeks later in the same manner as the Test Example 1. In addition, the presence of itchiness of the skin surface and skin condition were  
20 evaluated after 4 weeks. Furthermore, the evaluation scores in the above Test Example 1 were used for evaluation.

Those results are shown in Table 5 below.

[Table 5]

Patient	Start of treatment	After 3 days	After 1 week	After 2 weeks	After 3 weeks	After 4 weeks	Overall evaluation	
							Itchiness	Skin surface
P	5	4	3	1	1	1	none	Normal
Q	5	4	3	3	1	1	none	Normal
R	5	5	3	2	1	1	none	Normal
S	5	4	2	1	1	1	none	Normal
T	5	4	2	1	1	1	none	Normal

25 As is clear from the above results, the external preparation of the present invention was observed to

demonstrate improvement of dermatitis symptoms in 3-7 days after the start of application during treatment of atopic dermatitis, and the skin was no different from complete normal skin 3-4 weeks after the start of application.

Furthermore, there was no irritation of the skin by the preparation during application. In addition, there were also no adverse side effects such as rebound observed, which are observed with steroid-based external preparations, even after administration was discontinued.

#### (Text Example 5) Treatment of Atopic Dermatitis

The therapeutic effect of the cream produced in the above Example 21 was examined by applying to patients with atopic dermatitis.

The cream was applied to the target patients indicated below.

Target patient U: 40-year-old female suffering from atopic dermatitis

Target patient V: 38-year-old female suffering from atopic dermatitis

Target patient W: 55-year-old female suffering from atopic dermatitis

#### Method:

For target patients V and W, the cream for external use produced in Example 21 was applied twice a day for 4 consecutive weeks to the face that exhibited prominent atopic dermatitis, and the status of inflammation was observed.

For target patient U, a cream for external use containing only metronidazole was applied twice a day for 4 consecutive weeks to the face that exhibited prominent atopic dermatitis, and the status of inflammation was observed.

Therapeutic effects were evaluated by scoring rash, eczema and other dermatitis symptoms at the start of treatment along with the status of healing over 3 days, 1 week, 2 weeks, 3 weeks and 4 weeks later in the same manner as the Test Example 1. In addition, the presence of itchiness of the skin surface and skin condition were

evaluated after 4 weeks.

Furthermore, evaluation scores were determined in the manner indicated below.

Skin Condition:

- 5 5: Prominent rash, eczema and other dermatitis symptoms while also suffering from pain.
- 4: Prominent rash, eczema and other dermatitis symptoms, but not to the extent of grade 5.
- 10 3: Rash, eczema and other dermatitis symptoms able to be confirmed, but not to the extent of grade 4.
- 2: Rash, eczema and other dermatitis symptoms only able to be confirmed slightly, and not that much different from normal skin.
- 15 1: Absence of rash, eczema and other dermatitis symptoms, and appearance of normal skin.

Skin Itchiness:

- 3: Prominent itchiness, skin is scratched subconsciously.
- 2: Occasional itchiness, and scratching able to be controlled.
- 20 1: No itchiness whatsoever.

Those results are summarized in Table 6 below.

In Table 6, S1 refers to the score for skin condition, while S2 refers to the score for itchiness.

[Table 6]

Patient	Start of treatment	3 days after	1 week after	2 weeks after	3 weeks after	4 weeks after	Overall evaluation
	S1:S2	S1:S2	S1:S2	S1:S2	S1:S2	S1:S2	S1:S2
U	5:3	5:3	3:3	2:2	2:2	1:1	1:1
V	5:3	4:2	3:1	2:1	2:1	1:1	1:1
W	5:3	4:1	3:1	1:1	1:1	-	1:1

As indicated above, although patients U, V and W exhibited skin conditions no different from those of healthy individuals after 4 weeks, patients V and W, who applied the cream for external use of Example 21, which is a compound

preparation, no longer exhibited itchiness sooner than patient U, who applied an external preparation containing metronidazole alone. In addition, improvement of the skin was also faster in patients V and W. Furthermore, since symptoms of atopic dermatitis were no longer able to be confirmed for patient W in week 3, application was discontinued in week 3 at that patient's request.

(Text Example 6) Treatment of Atopic Dermatitis

The therapeutic effect of the creams produced in the above Examples 22 and 51 were examined by applying to patients with atopic dermatitis.

Test Method:

The cream for external use produced in the Example 22 was applied twice a day for 4 consecutive weeks to the right arm of target patient U of Test Example 5 having atopic dermatitis, and the status of inflammation was observed. In addition, the cream for external use produced in Example 51 was applied twice a day for 4 consecutive weeks to the left arm of target patient U of the Test Example 5 having atopic dermatitis, and the status of inflammation was observed.

Therapeutic effects were evaluated by scoring rash, eczema and other dermatitis symptoms at the start of treatment along with the status of healing over 3 days, 1 week, 2 weeks, 3 weeks and 4 weeks later. In addition, the presence of itchiness of the skin surface and skin condition were evaluated after 4 weeks.

Furthermore, the same evaluation scores as used in Test Example 5 were used for evaluation.

Those results are summarized in Table 7.

In Table 7, S1 refers to the score for skin condition, while S2 refers to the score for itchiness.

[Table 7]

Target	Start of treat- ment S1:S2	After 3 days S1:S2	After 1 week S1:S2	After 2 weeks S1:S2	After 3 weeks S1:S2	After 4 weeks S1:S2	Overall evalua- tion S1:S2
Right arm	4:3	3:1	2:1	2:1	1:1	1:1	1:1
Left arm	4:3	4:3	3:2	2:2	2:1	1:1	1:1

As is indicated above, the skin conditions of the left and right arms exhibiting the same symptoms in the same patient U improved after 4 weeks. Itchiness disappeared more quickly and the skin improved more rapidly on the right arm, to which was applied the cream for external use produced in Example 22 in the form of a compound preparation, than on the left arm, to which was applied the cream for external use of Example 51 in the form of an external preparation of metronidazole alone.

#### (Test Example 7) Treatment of Atopic Dermatitis

The therapeutic effect of the cream produced in the above Example 23 was examined by applying to patients with atopic dermatitis.

The cream was applied to the target patients indicated below.

Target patient X: 30-year-old female suffering from atopic dermatitis

Target patient Y: 28-year-old female suffering from atopic dermatitis

Target patient Z: 26-year-old female suffering from atopic dermatitis

Target patient a: 50-year-old female suffering from atopic dermatitis on the head

#### Method:

For target patient X, the cream for external use containing only tinidazole was applied twice a day for 4 consecutive weeks to the face that exhibited prominent atopic dermatitis, and the status of inflammation was observed.

For target patients Y and Z, a cream for external use produced in the Example 23 was applied twice a day for 4 consecutive weeks to the face that exhibited prominent atopic dermatitis, and the status of inflammation was observed.

5 For target patient a, the gel produced in the Example 25 was applied 2-3 times a day until symptoms improved, and those effects were observed.

10 Therapeutic effects were evaluated by scoring rash, eczema and other dermatitis symptoms at the start of treatment along with the status of healing over 3 days, 1 week, 2 weeks, 3 weeks and 4 weeks later in the same manner as the Test Example 1. In addition, the presence of itchiness of the skin surface and skin condition were evaluated after 4 weeks.

15 Furthermore, the same evaluation scores as used in the Test Example 5 were used for evaluation.

Those results are summarized in Table 8.

In Table 8, S1 refers to the score for skin condition, while S2 refers to the score for itchiness.

20 [Table 8]

Patient	Start of treat- ment	After 3 days	After 1 week	After 2 weeks	After 3 weeks	After 4 weeks	Overall evaluation
	S1:S2	S1:S2	S1:S2	S1:S2	S1:S2	S1:S2	S1:S2
X	5:3	5:3	4:2	3:2	2:1	2:1	2:1
Y	5:3	4:1	3:1	2:1	1:1	1:1	1:1
Z	4:3	3:1	2:1	1:1	--	--	1:1
a	4:3	3:1	3:1	2:1	2:1	1:1	1:1

25 As is indicated above, although target patients X, Y, Z and a exhibited skin conditions not different from those of healthy individuals after 4 weeks, itchiness disappeared more quickly and the skin improved more rapidly in patients Y, Z and a, who applied compound preparations in form of the cream for external use of the Example 23 or the gel of the Example 25, than patient Y, who applied an external preparation

containing tinidazole alone. Furthermore, since symptoms of atopic dermatitis were unable to be confirmed for patient Z in week 2, application was discontinued in week 2 at that patient's request.

5

(Test Example 8) Treatment of Atopic Dermatitis

The therapeutic effect of an external preparation of the present invention was examined by applying to an actual patient with atopic dermatitis. The external preparation was applied to the target patients indicated below.

Target patient b: 40-year-old male suffering from atopic dermatitis

Method:

The cream for external use of the Example 11 was applied twice a day for 4 consecutive weeks to the left arm of target patient b having atopic dermatitis, and the status of inflammation was observed.

In addition, the cream for external use of the Example 24 was applied twice a day for 4 consecutive weeks to the right arm of the same target patient b having atopic dermatitis, and the status of inflammation was observed.

Therapeutic effects were evaluated by scoring rash, eczema and other dermatitis symptoms at the start of treatment along with the status of healing over 3 days, 1 week, 2 weeks, 3 weeks and 4 weeks later. In addition, the presence of itchiness of the skin surface and skin condition were evaluated after 4 weeks.

Furthermore, the same evaluation scores as used in Test Example 5 were used for evaluation.

Those results are summarized in Table 9.

In Table 9, S1 refers to the score for skin condition, while S2 refers to the score for itchiness.

35

[Table 9]

Target	Start of treat- ment S1:S2	After 3 days S1:S2	After 1 week S1:S2	After 2 weeks S1:S2	After 3 weeks S1:S2	After 4 weeks S1:S2	Overall evaluation S1:S2
Right arm	5:3	5:1	4:1	2:1	2:1	2:1	1:1
Left arm	5:3	5:3	4:2	4:2	3:1	3:1	3:1

As is indicated above, the skin conditions of the left and right arms exhibiting the same symptoms in the same patient b improved after 4 weeks. Itchiness disappeared more quickly and the skin improved more rapidly on the right arm, to which was applied the cream for external use of Example 24 in the form of a compound preparation, than on the left arm, to which was applied the cream for external use of Example 11 in the form of an external preparation of tinidazole alone.

(Test Example 9)

The therapeutic effects of external preparations produced in the examples were examined by applying to actual patients with eczema/rash and seborrheic dermatitis.

The external preparations were applied to the target patients indicated below.

Target patient c: 60-year-old female suffering from cosmetic rash.

Target patient d: 34-year-old male suffering from seborrheic dermatitis.

Target patient e: 45-year-old male suffering from insect bite (mite).

Target patient f: 57-year-old male suffering from tinea.

Target patient g: 30-year-old female suffering from acne.

Target patient h: 28-year-old male suffering from suppurative dermatitis on the head.

Target patient i: 25-year-old male suffering from dermatitis herpiformis (blisters) on the neck.

Target patient j: 45-year-old female suffering from

candidiasis between the fingers.

Target patient k: 63-year-old male suffering from dry eczema on the back.

5 Target patient l: 28-year-old male suffering from suppurative dermatitis on the head.

Target patient m: 63-year-old male suffering from boils and eczema on the neck.

Target patient n: 33-year-old male suffering from early herpes on the forehead.

10 Target patient o: 23-year-old female suffering from dry eczema on the lower limbs.

Method:

15 The cream for external use produced in the Example 22 was applied twice a day for 4 consecutive weeks to target patients c and d, and its effect was observed.

The ointment for external use produced in the Example 33 was applied twice a day until symptoms improved to target patient e, and its effect was observed.

20 The cream for external use produced in the Example 31 was applied twice a day for 4 consecutive weeks to target patient f, and its effect was observed.

The cream for external use produced in the Example 30 was applied twice a day until symptoms improved to target patient g, and its effect was observed.

25 The gel produced in the Example 39 was applied twice or three times a day until symptoms improved to target patient h, and its effect was observed.

30 The cream for external use produced in the Example 32 was applied twice or three times a day until symptoms improved to target patient i, and its effect was observed.

The ointment for external use produced in the Example 35 was applied twice a day until symptoms improved to target patient j, and its effect was observed.

35 The lotion for external use produced in the Example 36 was applied twice or three times a day for 4 consecutive weeks to target patient k, and its effect was observed.

The patch produced in the Example 37 was applied twice

or three times a day for 3 consecutive weeks to target patient 1, and its effect was observed.

The plaster produced in Example 38 was applied twice or three times a day for 3 consecutive weeks to target patient m, and its effect was observed.

The cream for external use produced in the Example 21 was applied twice or forth a day until symptoms improved to target patient n, and its effect was observed.

The ointment for external use produced in the Example 34 was applied twice or three times a day until symptoms improved to target patient o, and its effect was observed.

Therapeutic effects were evaluated by scoring rash, eczema and other dermatitis symptoms at the start of treatment along with the status of healing over 3 days, 1 week, 2 weeks, 3 weeks and 4 weeks later. In addition, the presence of itchiness of the skin surface and skin condition were evaluated after 4 weeks.

Furthermore, in addition to using the same evaluation scores as used in Test Example 5 for evaluation, pain was also evaluated using the scores indicated below.

Pain Status:

3: Stinging pain

2: Pain not noticed unless affected area touched

1: No pain even if affected area touched

Those results are summarized in Table 10 below.

In Table 10, S1 refers to the score for skin condition, S2 refers to the score for itchiness, and S3 refers to the score for pain.

[Table 10]

Patient	Evalua- tion	Start of treat- ment	After 3 days	After 1 week	After 2 weeks	After 3 weeks	After 4 weeks	Overall evalua- tion
c	S1:S2	5:3	5:2	4:1	3:1	2:1	2:1	2:1
d	S1:S2	4:3	3:2	2:1	2:1	1:1	1:1	1:1

e	S1:S2	3:3	2:1	1:1	--	--	--	1:1
f	S1:S2	5:3	4:2	3:2	3:1	2:2	2:1	2:2
g	S1:S2	4:3	3:1	2:1	2:1	1:1	1:1	1:1
h	S1:S3	4:3	2:1	2:1	1:1	--	--	1:1
i	S1:S2	4:3	2:1	1:1	1:1	--	--	1:1
j	S1:S3	3:3	2:1	1:1	--	--	--	1:1
k	S1:S3	4:2	3:2	1:1	1:1	1:1	--	1:1
l	S1:S2	4:3	4:3	3:1	2:1	2:1	--	2:1
m	S1:S3	3:3	3:3	2:2	2:1	2:1	--	2:1
n	S1:S2	3:3	1:1	--	--	--	--	1:1
o	S1:S2	4:3	3:1	2:1	1:1	1:1	--	1:1

As is indicated above, during treatment of various skin diseases, application of the external preparations of the present invention resulted in improvement in symptoms being observed 3-7 days after the start of treatment, and the skin was no different from normal skin after 3-4 weeks. Although the skin of patient c became keloid due to the adverse side effects of using steroids for about six months, dermatitis had been relieved. Although patient f was not completely healed after 4 weeks due to having suffered from tinea over the long period of roughly 40 years, improvement in skin condition was remarkable. Furthermore, there was no irritation of the skin by the preparation during application. In addition, there were also no adverse side effects such as rebound observed, which are observed with steroid-based external preparations, even after administration was discontinued.

(Test Example 10)

The therapeutic effects of external preparations produced in the examples were examined by applying to actual patients with eczema/rash and seborrheic dermatitis.

The external preparations were applied to the target patients indicated below.

Target patient p: 10-year-old boy suffering from

psoriasis vulgaris on the instep of the foot.

Target patient q: 10-year-old boy suffering from psoriasis vulgaris on the lower limb.

5 Target patient r: 45-year-old male suffering from suppurative dermatitis caused by an insect bite.

Target patient s: 50-year-old female suffering from erythroderma on the face.

Target patient t: 20-year-old female suffering from acne.

10 Target patient u: 23-year-old female suffering from eczema on the upper arm

Target patient v: 50-year-old female suffering from atopic dermatitis on the head.

Target patient w: 63-year-old female suffering from tinea on the toes of the feet.

15 Target patient x: 65-year-old male suffering from a tumor on the neck (having both odor and pain).

Method :

20 The cream for external use produced in the Example 44 was applied twice a day for 4 consecutive weeks to target patients p and q, and its effect was observed.

The ointment for external use produced in the Example 46 was applied twice a day until symptoms improved to target patient r, and its effect was observed.

25 The cream for external use produced in the Example 41 was applied twice a day for 4 consecutive weeks to target patient s, and its effect was observed.

The cream for external use produced in the Example 40 was applied twice a day until symptoms improved to target patient t, and its effect was observed.

30 The cream for external use produced in the Example 42 was applied twice or three times a day until symptoms improved to target patient u, and its effect was observed.

35 The gel produced in the Example 25 was applied twice or three times a day until symptoms improved to target patient v, and its effect was observed.

The lotion produced in the Example 47 was applied twice or three times a day until symptoms improved to target

patient w, and its effect was observed.

The cream for external use produced in the Example 50 was applied twice or three times a day for 4 consecutive weeks to target patient x, and its effect was observed.

5 Therapeutic effects were evaluated by scoring rash, eczema and other dermatitis symptoms at the start of treatment along with the status of healing over 3 days, 1 week, 2 weeks, 3 weeks and 4 weeks later. In addition, the presence of itchiness of the skin surface and skin condition  
10 were evaluated after 4 weeks.

Furthermore, the same evaluation scores as used in the Test Example 10 were used for evaluation, pain was also evaluated using the scores indicated below.

Those results are summarized in Table 11 below.

15 In Table 11, S1 refers to the score for skin condition, S2 refers to the score for itchiness, and S3 refers to the score for pain.

[Table 11]

Patient	Evalu- ation	Start of treat- ment	After 3 days	After 1 week	After 2 weeks	After 3 weeks	After 4 weeks	Overall evalua- tion
p	S1:S2	4:3	3:2	2:1	2:1	1:1	1:1	1:1
q	S1:S2	4:3	2:1	2:1	2:1	1:1	1:1	1:1
r	S1:S3	4:3	2:1	2:1	1:1	1:1	-:-	1:1
s	S1:S2	5:2	4:1	3:1	2:1	2:1	2:1	2:1
t	S1:S2	3:3	1:1	1:1	-:-	-:-	-:-	1:1
u	S1:S2	4:3	2:1	2:1	1:1	-:-	-:-	1:1
v	S1:S2	4:3	3:1	3:1	2:1	2:1	1:1	1:1
w	S1:S2	4:3	2:1	2:1	2:1	1:1	1:1	1:1
x	S1:S3	4:3	4:3	3:2	3:2	2:1	2:1	2:1

20 As is indicated above, during treatment of various skin diseases, application of the creams for external use of the present invention resulted in improvement in symptoms being observed 3-7 days after the start of treatment, and the skin was no different from normal skin after 3-4 weeks. In

addition, although patient v noticed partial hair loss on the head, the hair began to grow somewhat starting in week 3.

Furthermore, there was no irritation of the skin by the preparation during application. In addition, there were also  
5 no adverse side effects such as rebound observed, which are observed with steroid-based external preparations, even after administration was discontinued.

(Test Example 11)

10 The therapeutic effects of external preparations of the present invention were examined by applying to actual patients with dermatitis and hircus.

The external preparations were applied to the target patients indicated below.

15 Target patient y (right): Right arm of a 33-year-old male suffering from hircus

Target patient y (left): Left arm of a 33-year-old male suffering from hircus

Method:

20 Target patient y (right): Cream for external use of the Example 52

Target patient y (left): Cream for external use of the Example 53

The above creams for external use were each applied  
25 twice a day to site affected by hircus, and progress was observed.

Therapeutic effects were evaluated by scoring odor and other symptoms at the start of treatment along with the status of healing over 3 days, 1 week, 2 weeks and 3 weeks  
30 later.

Furthermore, evaluation scores were determined in the manner indicated below.

Skin Soiling:

4: Soiled

35 3: Somewhat soiled

2: Almost clean

1: Clean

Odor:

4: Strong odor

3: Slight odor

2: Hardly any odor

5 1: No odor

Those results are summarized in Table 12 below.

In Table 12, S4 refers to the score for skin soiling,  
and S5 refers to the score for odor.

[Table 12]

Target	Start of treatment	After 3 days	After 1 week	After 2 weeks	After 3 weeks
	S4:S5	S4:S5	S4:S5	S4:S5	S4:S5
Right arm	4:4	4:4	3:3	3:1	1:1
Left arm	4:4	4:4	3:3	3:1	1:1

As indicated above, hircus on the right and left arms of target patient y was alleviated after 7-14 days, and was completely healed after 14-21 days.

Furthermore, there was no irritation or abnormalities of the skin of target patient y during application.

(Test Example 12) Treatment of Hircus

The therapeutic effects of external preparations of the present invention were examined by applying to actual patients with dermatitis and hircus.

The external preparations were applied to the target patients indicated below.

Target patient z: 33-year-old male suffering from hircus

Target patient Aa: 33-year-old male suffering from hircus

Method:

Target patient z: Cream for external use of the Example 22

Target patient Aa: Cream for external use of the Example 24

The above creams for external use were each applied twice a day to site affected by hircus, and progress was observed.

5 Therapeutic effects were evaluated by scoring odor and other symptoms at the start of treatment along with the status of healing over 3 days, 1 week, 2 weeks and 3 weeks later.

Furthermore, the same evaluation scores used in the Test Examples 10 and 12 were used for evaluation.

10 Those results are summarized in Table 13 below.

In Table 13, S1 refers to the score for skin condition, and S5 refers to the score for odor.

[Table 13]

Target patient	Evaluation	Start of treatment	After 3 days	After 1 week	After 2 weeks	After 3 weeks	After 4 weeks	Overall evaluation
z	S1:S5	4:3	2:1	1:1	1:1	-:-	-:-	1:1
Aa	S1:S5	3:3	2:1	1:1	-:-	-:-	-:-	1:1

15 As indicated above, the external preparations of the present invention improved the skin condition and odor associated with hircus in a short period of time.

(Test Example 13) Treatment of Hircus

20 The therapeutic effects of external preparations of the present invention were examined by applying to actual patients with dermatitis and hircus.

Target patient Ab: Right arm of 27-year-old male suffering from hircus

25 Target patient Ac: Left arm of 27-year-old male suffering from hircus

Target patient Ad: Right arm of 44-year-old male suffering from hircus

30 Target patient Ae: Left arm of 44-year-old male suffering from hircus

Target patient Af: Right arm of 23-year-old female  
suffering from hircus

Target patient Ag: Left arm of 23-year-old female  
suffering from circus

5 Method :

Target patient Ab: Cream for external use of the Example  
54

Target patient Ac: Cream for external use of the Example  
55

10 Target patient Ad: Cream for external use of the Example  
56

Target patient Ae: Cream for external use of the Example  
57

15 Target patient Af: Cream for external use of the Example  
58

Target patient Ag: Cream for external use of the Eample  
59

20 The above creams for external use were each applied  
twice a day to site affected by hircus, and progress was  
observed.

Therapeutic effects were evaluated by scoring odor and  
other symptoms at the start of treatment along with the  
status of healing in the course of time.

25 Furthermore, the same evaluation scores used in the Test  
Example 11 were used for evaluation.

Those results are summarized in Table 14 below.

In Table 14, S4 refers to the score for skin soiling,  
and S5 refers to the score for odor.

30 [Table 14]

Target patient	Evalua- tion	Start of treatment	After 1 day	After 3 days	After 5 days	After 7 days	After 10 days
Ab	S4:S5	4:4	4:4	3:2	2:1	2:1	1:1
Ac	S4:S5	4:4	4:3	3:2	3:1	2:1	1:1
Ad	S4:S5	4:4	4:3	3:2	3:1	2:1	1:1
Ae	S4:S5	4:3	4:3	3:1	3:1	1:1	1:1

Af	S4:S5	4:4	3:3	2:3	2:1	2:1	1:1
Ag	S4:S5	4:4	4:3	3:3	3:1	2:1	2:1

---

As indicated above, odor associated with hircus was alleviated after 3-5 days for all of the target patients, and both hircus and skin condition improved after 7-10 days.

Furthermore, there was no irritation or abnormalities of the skin of any of the subjects during application.

Similar results were obtained against hircus for the external preparations of the Example 25 and the Examples 60-64, with odor improving after about 3-5 days, and both odor and skin condition improving after 7-10 days. In addition, lotions and gels were easy to use.

Furthermore, since it was possible that hircus may have been alleviated due to the effects of the base alone, a placebo was applied twice a day for about 2 consecutive weeks. This had no effect, however.

#### (Test Example 14) Treatment of Odor

The therapeutic effects of external preparations of the present invention were examined by applying to actual patients with foot odor.

The external preparations were applied to the target patients indicated below.

Target patient Ah: Right foot of 24-year-old male having foot odor starting below the ankle (at the location from where foot odor is usually thought to emit).

Target patient Ai: Left foot of 24-year-old male having foot odor starting below the ankle (at the location from where foot odor is usually thought to emit).

#### Method:

Target patient Ah: Lotion for external use of the Example 63

Target patient Ai: Placebo lotion for external use from which the active ingredient of the Example 63 had been removed.

When the lotion for external use containing the active

ingredient was applied to the right foot, foot odor disappeared in about 4-5 hours. However, when the placebo lotion was applied to the left foot, foot odor persisted even after about 4-5 hours.

5

(Test Example 15)

The therapeutic effects of external preparations of the present invention were examined by applying to actual patients with dermatitis and psoriasis.

10 The external preparations were applied to the target patients indicated below.

Target patient Aj: Right foot of 43-year-old male suffering from psoriasis

15 Target patient Ak: Left foot of 43-year-old male suffering from psoriasis

Target patient Al: Right foot of 40-year-old male suffering from psoriasis

Target patient Am: Left foot of 40-year-old male suffering from psoriasis

20 Target patient An: Right foot of 38-year-old female suffering from psoriasis

Target patient Ao: Left foot of 38-year-old female suffering from psoriasis

25 Target patient Ap: Right foot of 49-year-old female suffering from psoriasis

Target patient Aq: Left foot of 49-year-old female suffering from psoriasis

Method :

30 Target patient Aj: Cream for external use of the Example 52

Target patient Ak: Cream for external use of the Example 53

Target patient Al: Cream for external use of the Example 65

35 Target patient Am: Cream for external use of the Example 66

Target patient An: Cream for external use of the Example

130

Target patient Ao: Cream for external use of the Example

131

Target patient Ap: Cream for external use of the Example

67

Target patient Aq: Cream for external use of the Example

68

The above creams for external use were each applied three times a day to the site affected by psoriasis, and progress was observed.

Furthermore, evaluation scores were determined in the manner indicated below.

Evaluation:

5: Worse than at the start of treatment

4: No change from the start of treatment

3: Some improvement

2: Remarkable improvement

1: No different from normal skin

Those results are summarized in Table 15 below.

[Table 15]

Target patient	After 3 days	After 7 days	After 21 days	After 1 month	After 2 months	After 3 months
Aj	4	3	3	2	2	1
Ak	4	3	3	2	2	1
Al	4	2	2	2	1	(1)
Am	4	2	2	2	1	(1)
An	4	2	2	2	1	1
Ao	4	2	2	1	1	1
Ap	4	2	2	1	1	(1)
Aq	4	2	2	1	1	(1)

As indicated above, symptoms of psoriasis exhibited significant change after 7-21 days in all of the target patients, and psoriasis was completely healed after 1-2 months.

5 Although application was discontinued after 2 months for target patients Al, Am, Ap and Aq, there was no recurrence as of 1 month later. In addition, there were no adverse side effects and so forth observed in any of the target patients.

10 (Test Example 16) Treatment of Psoriasis

Target patient Ar: Right foot of 38-year-old male suffering from psoriasis

Target patient As: Left foot of 38-year-old male suffering from psoriasis

15 Method:

Target patient Ar: Cream for external use of the Example 65

Target patient As: Commercially available Bonalfa ointment (Teijin Co.,Ltd.) (ingredient: tacalcitol)

20 The above creams for external use were each applied twice a day to the site affected by psoriasis, and their respective therapeutic effects were observed in the course of time.

Furthermore, the same evaluation scores used in the Test Example 15 were used for evaluation.

Those results are summarized in Table 16 below.

[Table 16]

Target patient	After 3 days	After 7 days	After 21 days	After 1 month	After 2 months	After 3 months
Ar	4	2	2	1	-	-
As	4	4	3	2	-	-

30 As indicated above, skin condition clearly improved for target patient As, namely with application of the

metronidazole external preparation. There were no adverse side effects observed for either preparation.

(Test Example 17)

5 Target patient At: Head of 33-year-old male suffering from psoriasis

Target patient Au: Right elbow of 38-year-old male suffering from psoriasis

10 Target patient Av: Left elbow of 38-year-old male suffering from psoriasis

Method:

Target patient At: Lotion of the Example 72

Target patient Au: Composite preparation for external use of the Example 73

15 Target patient Av: Cream for external use of the Example 65

20 The above creams for external use were each applied twice a day to the site affected by psoriasis, and their respective therapeutic effects were observed in the course of time.

Furthermore, the same evaluation scores used in Test Example 15 were used for evaluation.

Those results are summarized in Table 17 below.

[Table 17]

Target patient	After 3 days	After 7 days	After 21 days	After 1 month	After 2 months	After 3 months
At	4	2	1	1	-	-
Au	4	2	2	1	1	-
Av	4	3	2	2	1	-

25

As indicated above, remarkable improvement was observed after 7 to 21 days. With respect to a comparison of target patients Au and Av, a compound metronidazole preparation (2%) was applied to the right foot, and a preparation containing metronidazole alone (2%) was applied to the left foot. A comparison revealed the compound preparation to be more

30

effective. In addition, there were no adverse side effects observed for any of the preparations.

(Test Example 18) Treatment of Psoriasis

5        Testing was performed on target patients Aw, Ax, Ay, Az, Ba and Bb in the same manner as the Test Example 15.

Method:

      Target patients Aw and Ax: Ointment for external use of the Example 69

10       Target patients Ay and Az: Ointment for external use of the Example 70

      Target patient Ba: Lotion of the Example 72

      Target patient Bb: Composite preparation of the Example 74

15       The external preparations of the present invention demonstrated remarkable effects in all of these target patients as well, and four of the target patients were completely healed within 1 month, while the remaining two target patients were completely healed within 1-3 months.

20       (Test Example 19) Treatment of Psoriasis

      Target patient Bc: Right foot of 70-year-old patient suffering from psoriasis

      Target patient Bd: Left foot of 70-year-old patient suffering from psoriasis

25       Method:

      Target patient Bc: cream for external use of the Example 73

30       Target patient Bd: cream for external use prepared in the same manner as the Example 73 but without adding metronidazole (preparation containing Tranilast only)

      The above creams for external use were each applied twice a day to the site affected by psoriasis, and their respective therapeutic effects were observed in the course of time.

35       Furthermore, the same evaluation scores used in the Test Example 15 were used for evaluation.

Those results are summarized in Table 18 below.

[Table 18]

Target patient	After 3 days	After 7 days	After 21 days	After 1 month	After 2 months	After 3 months
Bc	4	2	2	2	1	1
Bd	4	4	4	4	3	4

5 As indicated above, there were no effects observed at all in the case of applying Tranilast at 0.1%.

(Test Example 20) Use for Scars and Blotches

Target patients:

10 Target patient Be: Right arm of 40-year-old male having a scar

Target patient Bf: Right arm of 40-year-old male having a scar

15 Target patient Bg: Face of 38-year-old female having blotches

Target patient Bh: Face of 60-year-old male having blotches

Target patient Bi: Right foot of 27-year-old patient having a scar

20 Target patient Bj: Left foot of 27-year-old patient having a scar

Method :

Be: External preparation of the Example 75

Bf: External preparation of the Example 76

25 Bg: External preparation of the Example 78

Bh: External preparation of the Example 77

Bi: External preparation of the Example 77

Bj: External preparation prepared in the same manner as the Example 75 but without adding metronidazole

30 The above external preparations were each applied three times a day (twice a day for target patients Bi and Bj), and

the progress was observed.

Furthermore, evaluation scores were determined in the manner indicated below.

Evaluation:

- 5    3:    Status prior to administration or no change  
     2:    Improvement over previous condition  
     1:    Clear improvement as compared with grade 2

Those results are summarized in Table 19 below.

10   [Table 19]

Target patient	Start of treatment	After 2 weeks	After 3 weeks	After 1 month	After 2 months
Be	3	3	3	2	2
Bf	3	3	2	2	1
Bg	3	3	2	1	1
Bh	3	3	2	2	1
Bi	3	3	2	1	1
Bj	3	3	3	3	3

As indicated by the above results, skin condition improved, and there are no particular adverse side effects observed. In addition, the skin had more luster and was smoother than before application. There were no particular changes in target patient Bj to which only a base was applied.

15

(Test Example 21)

Target patients:

20        Target patient Bk: Right hand of 34-year-old male on which the skin has been damaged by a burn

         Target patient Bl: Right finger of 33-year-old male on which there is a wound from a cut

         Target patient Bm: Right hand of a 34-year-old male on

25

which the skin has been damaged following the removal of a wart

Target patient Bn: Right foot of 12-year-old boy having a scrape wound

5 Target patient Bo: Left foot of 12-year-old boy having a scrape wound

Target patient Bp: Face of 5-year-old infant on which the skin has been damaged by a scratch

10 Target patient Bq: Right arm of 5-year-old infant on which the skin has been damaged by a scratch

Target patient Br: Left arm of 5-year-old infant on which the skin has been damaged by a scratch

Method:

15 Bk: External preparation of the Example 79

B1: External preparation of the Example 81

Bm: External preparation of the Example 82

Bn: External preparation of the Example 79

Bo: External preparation of the Example 81

Bp: External preparation of the Example 80

20 Bq: External preparation of the Example 82

Br: External preparation prepared in the same manner as the Example 79 but without adding metronidazole

25 The above external preparations were each applied three times a day (twice a day for target patients Bq and Br), and the progress was observed.

Furthermore, evaluation scores were determined in the manner indicated below.

Evaluation:

30 3: Status prior to administration or no change

2: Improvement over previous condition

1: Clear improvement as compared with grade 2

Those results are summarized in Table 20 below.

35 In Table 20, S6 refers to the score for skin condition, S7 refers to the score for pain, and S8 refers to the score for irritation.

[Table 20]

Target patient	Evaluation	Start of treatment	After 1 day	After 3 days	After 1 week	After 2 weeks
Bk	S6:S7	3:3	3:2	1:1	1:1	--
Bl	S6:S7	3:3	3:2	1:1	1:1	--
Bm	S6:S7	3:3	3:2	2:2	1:2	1:1
Bn	S6:S7	3:3	3:3	2:1	1:1	--
Bo	S6:S7	3:3	3:2	2:1	1:1	--
Bp	S6:S8	3:3	3:2	1:1	1:1	1:1
Bq	S6:S8	3:3	2:2	2:1	1:1	1:1
Br	S6:S8	3:3	3:3	3:3	3:3	--

As indicated by the above results, skin condition improved, and there was no occurrence of adverse side effects. These preparations were particularly superior with respect to improving or eliminating pain quite rapidly. In addition, the skin also had more luster and was smoother than before application. Since there were no particular changes observed in target patient Br in which only a base was applied, application of base only was discontinued after 1 week at the request of the target patient and parents, and when the cream of the Example 82 was applied instead, the skin was nearly completely healed in 1-2 weeks.

(Test Example 22)

Target patients:

Target patient Bs: Right foot of 22-year-old female suffering from skin disease caused by weed ill

Target patient Bs: Right hand of 22-year-old female suffering from skin disease caused by weed ill

Target patient Bu: Two parts of face of 27-year-old male suffering from an insect bite

Target patient Bv: Right hand of 27-year-old male suffering from an insect bite

Target patient Bw: Right hand of 24-year-old female  
suffering from contact dermatitis

Target patient Bx: Right foot of 47-year-old male  
suffering from contact dermatitis

5 Target patient By: 28-year-old female suffering from  
dermatitis caused by detergent rash

Method:

Bs: External preparation of the Example 83

Bt: External preparation of the Example 85

10 Bu: External preparation of the Example 84

Bv: External preparation of the Example 86

Bw: External preparation of the Example 86

Bx: External preparation of the Example 86

By: External preparation of the Example 85

15 The above external preparations were each applied two to  
three times a day (twice a day for target patients Bu and Bv,  
whenever the hands were washed for target patients Bw and By),  
and the progress was observed.

20 Furthermore, evaluation scores were determined in the  
manner indicated below.

Evaluation:

3: Status prior to administration or no change

2: Improvement over previous condition

1: Clear improvement as compared with grade 2

25 Those results are summarized in Table 21 and Table 22  
below.

[Table 21]

Target patient	Start of treatment	After 1 hour	After 3 hours	After 6 hours	After 1 day	After 3 days
Bs	3	3	2	1	1	-
Bt	3	3	2	1	1	-
Bu	3	3	2	2	1	1
Bv	3	3	2	1	1	1

[Table 22]

Target	Start of treatment	After 3 days	After 7 days	After 2 wees	After 1 month	After 3 months
Bw	3	3	2	2	1	1
Bx	3	3	3	2	2	1
By	3	2	1	1	-	-

As indicated by the above results, skin condition improved, and there was no occurrence of adverse side effects.

5 These preparations were proved to eliminate itching discomfort and pain in a short period of time in particular. Although scars from the insect bites remained for target patients Bu and Bv, the scars disappeared about 1 week after the start of application. Although improvement similarly  
10 took some time for target patients Bw and Bx as well, itching was improved or eliminated in about 3-7 days after application.

(Test Example 23)

15 Target patients:

Target patient Bz: Back of a 78-year-old male suffering from dry pruritis

Target patient Ca: Back of 71-year-old male suffering from eczema through to a drug-induced side effect (anti-  
20 hypertension drug)

Target patient Cb: Back of 83-year-old male suffering from eczema

Target patient Cc: Both of arm of 83-year-old male suffering from eczema

25 Target patient Cd: Back of 68-year-old female suffering from dry pruritis

Target patient Ce: Face of 30-year-old female suffering from eczema caused by a cosmetics-induced side effect

30 Target patient Cf: Face of 40-year-old female suffering from eczema caused by a cosmetics-induced side effect

Method:

The external preparation of the Example 87 to target patients Bz, Ca, Cb, Cc and Ce, and the external preparation of the Example 88 to target patients Cd and Cf were each applied twice a day, and the progress was observed.

5 Furthermore, evaluation scores were determined in the manner indicated below.

Evaluation:

3: Status prior to administration or no change

2: Improvement over previous condition

10 1: Clear improvement as compared with grade 2

Those results are summarized in Table 23 below.

[Table 23]

Target patient	Start of treatment	After 3 days	After 1 week	After 2 weeks	After 1 month
Bz	3	1	1	1	-
Ca	3	2	2	1	1
Cb	3	2	1	1	-
Cc	3	2	1	1	-
Cd	3	1	1	1	1
Ce	3	3	2	2	1
Cf	3	3	2	2	1

15 As indicated by the above results, skin condition improved, and there was no occurrence of adverse side effects. Itching stopped in a few days, and skin condition improved each day. Although target patients Ce and Cf required about 1 month for skin condition to heal completely due to the particularly serious cosmetic rash exhibited by these target patients, itching disappeared after about 3 days.

20

(Test Example 24)

Target patients:

25 Target patient Cg: Right hand of 26-year-old male

suffering from chapped skin

Target patient Ch: Left hand of 26-year-old male  
suffering from chapped skin

Target patient Ci: Right foot of 26-year-old male  
5 suffering from chilblain

Target patient Cj: Left foot of 26-year-old male  
suffering from chilblain

Method:

Cg: External preparation of the Example 89

10 Ch: External preparation produced in the same manner as  
the Example 89 but without using metronidazole

Ci: External preparation of the Example 90

Cj: External preparation produced in the same manner as  
the Example 89 but without using metronidazole

15 The above external preparations were each applied twice  
a day for target patients Ci and Cj, whenever the hands were  
washed for target patients Cg and Ch, and the progress was  
observed.

20 Furthermore, evaluation scores were determined in the  
manner indicated below.

Evaluation:

3: Status prior to administration or no change

2: Improvement over previous condition

1: Clear improvement as compared with grade 2

25 Those results are summarized in Table 24 below.

[Table 24]

Target patient	Start of treatment	After 3 days	After 1 week	After 2 weeks	After 1 month
Cg	3	3	2	2	2
Ch	3	3	3	3	3
Ci	3	2	2	2	1
Cj	3	3	3	3	3

As indicated by the above results, skin condition improved, and there was no occurrence of adverse side effects. Although skin condition was not improved by the preparations that did not contain the active ingredient, in those preparations that did contain the active ingredient, clear improvement was observed. In particular, itching and discomfort were improved in about 1 week.

(Test Example 25)

Target patients:

Target patient Ck: Back of a 74-year-old male suffering from dry erythroderma

Target patient Cl: Arm of a 74-year-old male suffering from dry erythroderma

Target patient Cm: Back of an 80-year-old male suffering from pustular psoriasis erythroderma

Method:

The external preparation of the Example 91 was applied to target patients Ck and Cm, and the external preparation of the Example 92 was applied to target patient Cl twice a day, and the progress was observed.

Furthermore, evaluation scores were determined in the manner indicated below.

Evaluation:

- 5: Skin condition worse than before administration
- 4: Skin condition before administration or no change
- 3: Some improvement as compared with before administration
- 2: Greater improvement as compared with grade 3
- 1: Clear improvement as compared with grade 2

Those results are summarized in Table 25 below.

[Table 25]

Target patient	Start of treatment	After 3 days	After 1 week	After 2 weeks	After 1 month	After 2 months
Ck	4	4	3	3	2	2
Cl	4	4	3	3	2	2

Cm	4	4	3	3	2	2
----	---	---	---	---	---	---

---

As indicated by the above results, skin condition improved, and there was no occurrence of adverse side effects. Itching improved in about 1 week. More time was required due to the nature of these skin conditions in being difficult to heal completely.

(Test Example 26)

Target patients:

10 Target patient Cn: Right foot of 55-year-old male suffering from tinea

Target patient Co: Left foot of 55-year-old male suffering from tinea

15 Target patient Cp: Right hand of 46-year-old female suffering from nail tinea

Target patient Cq: Right hand of 38-year-old female suffering from nail tinea

Method:

20 The external preparation of the Example 93 was applied to target patient Cn twice a day, the external preparation of the Example 94 was applied to target patient Co twice a day, the external preparation of the Example 94 was applied to subject Cp three times a day, and the external preparation of the Example 93 was applied to target patient Cq three times a day, and the progress was observed.

Furthermore, evaluation scores were determined in the manner indicated below.

Evaluation:

- 5: Skin condition worse than before administration
- 30 4: Skin condition before administration or no change
- 3: Some improvement as compared with before administration
- 2: Greater improvement as compared with grade 3
- 1: Clear improvement as compared with grade 2

Those results are summarized in Table 26 below.

[Table 26]

Target patient	Start of treatment	After 1 week	After 2 weeks	After 3 weeks	After 1 month	After 2 months
Cn	4	4	3	3	2	2
Co	4	3	3	2	2	2
Cp	4	3	2	2	1	1
Cq	4	3	2	2	2	2

As indicated by the above results, skin condition improved, and there was no occurrence of adverse side effects.

5 (Test Example 27)

Target patients:

Target patient Cr: Left foot of 49-year-old male suffering from suppurative skin disease

10 Target patient Cs: Area around the mouth of 61-year-old male suffering from herpes

Target patient Ct: Forehead of 33-year-old male suffering from herpes

Target patient Cu: Left arm of 64-year-old female suffering from suppurative skin disease

15 Target patient Cv: Right hand of 56-year-old subject suffering from candidiasis

Target patient Cw: Both hands of 38-year-old female suffering from periunguitis

20 Target patient Cx: Back of 33-year-old male suffering from dermal pruritis

Method:

The external preparation of the Example 96 was applied to target patient Cr three times a day, the external preparation of the Example 95 was applied to target patient  
25 Cs three times a day, the external preparation of Example 95 was applied to target patient Ct twice a day, the external preparation of the Example 96 was applied to target patient Cu three times a day, the external preparation of the Example 96 was applied to target patient Cv three to four times a day,

the external preparation of Example 96 was applied to target patient Cw whenever the hands were washed, and the external preparation of Example 95 was applied to target patient Cx twice a day, and the progress was observed.

5 Furthermore, evaluation scores were determined in the manner indicated below.

Evaluation:

4: Condition before administration or no change

3: Some improvement as compared with before administration

10 2: Definite improvement

1: No different from normal skin

Those results are summarized in Tables 27 and 28 below.

[Table 27]

Target patient	Start of treatment	After 1 day	After 3 days	After 1 week	After 2 weeks	After 1 month
Cs	4	2	1	1	-	-
Ct	4	1	1	1	-	-
Cx	4	2	1	1	-	-

15 [Table 28]

Target patient	Start of treatment	After 1 week	After 2 weeks	After 3 weeks	After 1 month	After 2 months
Cr	4	3	3	2	2	2
Cu	4	3	3	3	2	2
Cv	4	3	3	3	2	2
Cw	4	3	3	2	2	1

As indicated by the above results, skin condition improved, and there was no occurrence of adverse side effects. In all target patients, itchiness, pain and discomfort  
20 disappeared prior to improvement of skin condition.

(Test Example 28)

Target patients:

Target patient Cx: Face of 62-year-old female on which blotches are present on normal skin.

- 5 Target patient Cy: Face of 38-year-old male having a scar (in the form of a knob) caused by the adverse side effects of steroids.

Target patient Cz: Foot of 5-year-old boy having a scar caused by the adverse side effects of steroids

10 Method:

The cream for external use of the Example 97 was applied to the target patients twice a day.

Furthermore, evaluation scores were determined in the manner indicated below.

15 Evaluation:

5: Blotches and scars, etc. can be clearly distinguished from other parts of the skin.

4: Blotches and scars, etc. can be clearly distinguished from other parts of the skin but not to the degree of grade 5.

20 3: Blotches and scars, etc. can be distinguished from other parts of the skin.

2: Blotches and scars, etc. are slightly visible, but are essentially no different from other parts of the skin.

25 1: No difference whatsoever from other parts of the skin

Those results are summarized in Table 29 below.

[Table 29]

Target patient	Start of treatment	After 1 week	After 2 weeks	After 3 weeks	After 4 weeks	After 2 months	After 3 months
Cx	4	4	4	3	3	2	2
Cy	5	5	5	5	5	3	2
Cz	3	3	2	1	1	-	-

As indicated above, the external preparation of the present invention clearly demonstrated remarkable improvement with respect to blotches as well as scars caused by the adverse side effects of steroids.

5

(Test Example 29)

Target patients:

Target patient Da: Face of 63-year-old female with normal skin having a dark complexion.

10

Target patient Db: Face of 65-year-old female with normal skin having a dark complexion.

Method:

15

The external preparation of Example 98 was applied twice a day to target patient Da, while the preparation of Example 99 was applied twice a day to target patient Db.

Furthermore, evaluation scores were determined in the manner indicated below.

Evaluation:

20

- 3: Present skin condition of the face that would generally be considered to be dark.
- 2: Somewhat lighter complexion not to the extent of grade 3.
- 1: Clearly lighter complexion as compared with grade 3.

Those results are summarized in Table 30 below.

25

[Table 30]

Target	Start of treatment	After 2 weeks	After 1 month	After 2 months	After 3 months
Da	3	3	2	1	1
Db	3	3	2	1	1

As indicated above, the external preparations of the present invention clearly resulted in remarkable improvement of pigment deposition.

30

(Test Example 30)

Target patients:

Target patient Dc: Right foot of 54-year-old male

suffering from folliculitis

Target patient Dd: Back of 75-year-old male suffering from drug rash

5 Target patient De: 60-year-old male suffering from a laceration caused by contusion

Target patient Df: Left hand of 32-year-old male having pain caused by a scrape wound

Target patient Dg: Left shoulder of 44-year-old male suffering from suppurative skin disease

10 Target patient Dh: 38-year-old male whose eyebrows have become thin due to the adverse side effects of external steroid preparations

15 Target patient Di: Both arms of 38-year-old male on which scars remain due to the adverse side effects of external steroid preparations

Method:

20 The external preparation of the Example 100 was applied twice a day to target patient Dc, the external preparation of the Example 101 was applied twice a day to target patient Dd, the external preparation of the Example 102 was applied three times a day to target patient De, the external preparation of the Example 103 was applied twice a day to target patient Df, the external preparation of the Example 104 was applied twice a day to target patient Dg, the external preparation of the Example 105 was applied 3-4 times a day to target patient Dh, and the external preparation of the Example 106 was applied twice a day to target patient Di, and the progress was observed.

30 Furthermore, the evaluation scores for target patients Dc through Di were determined in the manner indicated below.  
Evaluation:

- 5: Skin symptoms are extremely worse  
4: Skin symptoms are extremely worse but not to the degree of grade 5  
35 3: Skin symptoms are moderate  
2: Skin symptoms are hardly able to be detected  
1: Normal skin

Those results are summarized in Table 31 below.

[Table 31]

Target patient	Start of treatment	After 3 days	After 1 week	After 2 weeks	After 3 weeks	After 4 weeks
Dc	4	3	1	1	-	-
Dd	3	1	1	1	1	1
De	3	2	1	1	-	-
Df	3	1	-	-	-	-
Dg	4	3	2	2	-	-
Di	5	5	4	4	3	3

5 Furthermore, evaluation scores for target patient Dh were determined in the manner indicated below.

Evaluation:

5: Eyebrows and vellus hair completely absent

4: Eyebrows absent but vellus hair growing

10 3: Slight eyebrow growth

2: Eyebrows have grown back but still visually conspicuous

1: Eyebrows identical to those of normal persons

Those results are summarized in Table 32 below

15

[Table 32]

Target patient	Start of treatment	After 3 days	After 1 week	After 2 weeks	After 3 weeks	After 4 weeks
Dh	5	5	4	3	3	2

20 As is clear from the above, the external preparations of the present invention exhibited ameliorative effects in each of the target patients.

Furthermore, target patient Dd suffered from hypertension and exhibited eczema and itching caused by the side effects of administration of an anti-hypertension drug.

Since the drug rash and itching reappeared when use of the external cream preparation was discontinued, this target patient used the preparation for a long period of time.

In addition, target patients Dh and Di refer to the same individual. Although skin condition was evaluated as grade 3 after 4 weeks, it was upgraded to grade 2 one month later. Although his eyebrows were also evaluated as grade 2 after 3 months as compared with the start of application, the number of hairs was increasing a little at a time.

(Test Example 31) Effects on Skin Moistness and Smoothness  
Target patients:

Target patient Dj: Right side of the face of 62-year-old female with normal skin.

Target patient Dk: Left side of the face of 62-year-old female with normal skin.

Target patient Dl: Right side of the face of 69-year-old female with normal skin.

Target patient Dm: Left side of the face of 69-year-old female with normal skin.

Target patient Dn: Left arm of 62-year-old female with normal skin.

Target patient Do: Right arm of 62-year-old female with normal skin.

Method:

The external preparation of the Example 107 to target patient Dj, an external preparation produced in the same manner as the Example 107 without using metronidazole to target patient Dk, the external preparation of the Example 108 to target patient Dl, and an external preparation produced in the same manner as the Example 108 without using tinidazole to target patient Dm, were respectively applied twice a day, while the external preparation produced in the Example 109 to target patient Dn and the external preparation produced in the Example 110 to target patient Do were respectively applied three times a day. All of the external preparations were applied for 2 consecutive months, and the

progress of the target patients was observed. Moreover, progress was also observed after application was discontinued 2 months after the start of application.

Furthermore, the evaluation scores were determined in the manner indicated below.

Evaluation (evaluation of skin moistness and smoothness when waking up on the day after application):

5: Worse skin condition than before use

4: Skin condition before use or no change

10 3: Some improvement as compared with before use

2: Definite improvement as compared with before use

1: Extremely good skin condition

Those results are summarized in Tables 33 and 34 below.

[Table 33] (Progress During Use)

15

Target patient	Before of treatment	After 3 days	After 1 week	After 2 weeks	After 3 weeks	After 4 weeks	After 2 months
Dj	4	4	3	2	2	1	1
Dk	4	4	4	4	4	4	3
Dl	4	4	3	2	2	1	1
Dm	4	4	4	4	4	4	4
Dn	4	3	2	2	2	1	1
Do	4	3	2	2	1	1	1

[Table 34] (Progress After Discontinuation of Application)

Target patient	When discontinued	After 3 days	After 1 week	After 2 weeks	After 1 month
Dj	1	1	1	2	2
Dk	3	4	4	4	4

Dl	1	1	1	1	2
Dm	4	4	4	4	4
Dn	1	1	1	2	2
Do	1	1	1	1	2

As indicated by the above results, satisfactory skin condition continued for 7-14 days after discontinuing application. There were no adverse side effects and the preparations were highly safe, while preparations not containing the active ingredient were not effective.

(Test Example 32)

Target patient Dp: Left foot of 7-year-old boy having a scratch wound suffering from atopic dermatitis.

Target patient Dq: Right foot of 7-year-old boy having a scratch wound suffering from atopic dermatitis.

Target patient Dr: Left foot of 39-year-old male having a scratch wound suffering from atopic dermatitis.

Target patient Ds: Right foot of 39-year-old male having a scratch wound suffering from atopic dermatitis.

Method :

Dp: Preparation of Example 112 applied twice a day.

Dq: Preparation of Example 113 applied twice a day.

Dr: Preparation of Example 114 applied three times a day

Ds: Preparation of Example 115 applied three times a day

Furthermore, the evaluation scores were determined in the manner indicated below.

5: Prominent rash, eczema and other dermatitis symptoms, extreme itchiness, subconscious scratching of the skin.

4: Prominent rash, eczema and other dermatitis symptoms along with considerable itchiness.

3: Rash, eczema and other dermatitis symptoms able to be confirmed and some itchiness.

2: Rash, eczema and other dermatitis symptoms only able to be confirmed slightly and not much different from normal skin, with some itchiness but to the extent that

scratching can be controlled.

- 1: Absence of rash, eczema and other dermatitis symptoms, appearance of normal skin and no itchiness.

Those results are shown in Table 35 below.

5

[Table 35]

Target patient	Start of treatment	After 3 days	After 7 days	After 2 weeks	After 3 weeks	After 1 months
Dp	4	4	3	2	1	1
Dq	4	4	3	2	2	2
Dr	5	5	4	3	3	2
Ds	5	5	4	3	3	2

Although ketoconazole and isoconazole nitrate can normally not be administered to affected areas where wounds are present, by producing compound preparations with nitroimidazole derivative in the external preparations of the present invention, these preparations were able to be used at affected areas where wounds were present. Although there was no change in skin condition in target patients Cp and Cq after about 3 days, itchiness had nearly completely disappeared, and scratching stopped after about 5 days even when sleeping.

Target patients Cr and Cs had a history of atopic dermatitis going back roughly 20 years, and skin condition had worsened considerably due to the adverse side effects of steroid drugs. Itchiness disappeared nearly completely after about 5 days, and there was no scratching after about 7 days.

(Test Example 33)

Target patients:

Target patient Dt: Left side of the back (from the waist to the shoulder) of 72-year-old male suffering from dermal pruritis

Target patient Du: Left side of the back (same location

as above) of 69-year-old female suffering from dermal pruritis

Target patient Dv: Right side of the back (same location as above) of 72-year-old male suffering from dermal pruritis

5 Target patient Dw: Right side of the back (same location as above) of 69-year-old female suffering from dermal pruritis

Furthermore, the evaluation scores were determined in the manner indicated below.

10 Method:

An external preparation produced in the same manner as Example 29 without using metronidazole to target patient Dt, an external preparation produced in the same manner as the Example 29 by adding 10 g of crotamiton without using  
15 metronidazole for target patient Du, the external preparation of the Example 111 for target patient Dv, and the external preparation of the Example 29 for target patient Dw were respectively applied twice a day.

Evaluation:

20 5: Itchiness worse than at the start of application  
4: Same as the start of application or extremely itchy  
3: Some alleviation of itchiness  
2: Occasional itchiness but hardly noticeable  
1: No itchiness whatsoever

25 Those results are summarized in Table 36 below.

[Table 36]

Target patient	Start of treatment	After 3 days	After 1 week	After 2 weeks	After 3 weeks	After 1 month	Overall evaluation
Dt	4	4	3	2	2	1	Hardly any change
Du	4	4	3	2	2	2	Some improvement
Dv	5	5	4	3	3	2	Remarkable improvement

							ment after 3 days to 1 week Remarkable improve- ment after 2-3 days
Dw	5	5	4	3	3	2	

As indicated above, in comparison with the preparation containing only crotamiton, the external preparations of the present invention demonstrated remarkable improvement of dermal pruritis. However, when crotamiton was contained, effects appeared earlier.

(Test Example 34) Stability Test

The ointment for external use produced in the above the Example 1, the cream for external use produced in the Example 4, the ointment for external use produced in the Example 11, and the cream for external use produced in the Example 14 were stored at room temperature and 40°C followed by observation of changes in their appearance, pH, content and viscosity after 6 months. Those results are shown in Table 37 below.

In Table 37, "NC" means no change as compared with before storing.

[Table 37]

Exampler	Appearance		pH		Content		Viscosity	
	room tem- perature	40°C	room tem- perature	40°C	room tem- perature	40°C	room tem- perature	40°C
1	NC	NC	NC	NC	NC	NC	NC	NC
4	NC	NC	NC	NC	NC	NC	NC	NC
11	NC	NC	NC	NC	NC	NC	NC	NC
14	NC	NC	NC	NC	NC	NC	NC	NC

The external preparations of the present invention exhibited no changes in appearance or pH, and did not exhibit any significant changes in content or viscosity.

Thus, the external preparations provided by the present  
5 invention were clearly determined to be pharmacologically stable.